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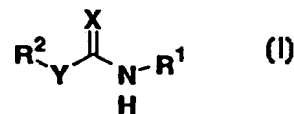
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(21) International Application Number: PCT/US98/02364 (22) International Filing Date: 6 February 1998 (06.02.98) (30) Priority Data: 60/038,811 21 February 1997 (21.02.97) US (71) Applicant: BRISTOL-MYERS SQUIBB COMPANY [US/US]; P.O. Box 4000, Princeton, NJ 08543-4000 (US). (72) Inventors: LLOYD, John; 1702 Yardley Road, Yardley, PA 19067 (US); ROVNYAK, George, C.; 10 W. Broad Street, Hopewell, NJ 08525 (US); STEIN, Philip, D.; 3 Harborton Ridge Drive, Pennington, NJ 08534 (US); AHMAD, Saleem; 2094 Chadi Drive, Wall, NJ 07719 (US); ATWAL, Karnail, S.; 92 Valley View Way, Newtown, PA 18940 (US); CAULFIELD, Thomas, J.; 4 Eric Court, Lawrenceville, NJ 08648 (US); POSS, Michael, A.; 15 Valerie Lane, Lawrenceville, NJ 08648 (US). (74) Agents: RODNEY, Burton et al.; Bristol Myers Squibb Company, P.O. Box 4000, Princeton, NJ 08543-4000 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>	

(54) Title: **BENZOIC ACID DERIVATIVES AND RELATED COMPOUNDS AS ANTIARRHYTHMIC AGENTS**

(57) Abstract

Benzoic acid derivatives of formula (I) where X is oxygen, sulfur, -NH, -NR¹, -N-CN, -N-OR¹ or -N-NO₂; Y is a single bond, -C=C-, or -NH; R¹ is alkyl, alkenyl, alkynyl, aryl, cycloalkyl, heterocyclo, or (heterocyclo)alkyl; and R² is aryl or heterocyclo. The compounds of formula (I) are useful in the treatment of arrhythmia.



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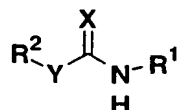
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BENZOIC ACID DERIVATIVES AND RELATED COMPOUNDS AS ANTIARRHYTHMIC AGENTS

5 **Brief Description of the Invention**

This invention is concerned with compounds of the formula

I



10 where -

X is oxygen, sulfur, -NH, -NR¹, -N-CN, -N-OR¹ or
-N-NO₂;

Y is a single bond, -C=C-, or -NH;

R¹ is alkyl, alkenyl, alkynyl, aryl, cycloalkyl, heterocyclo, or
15 (heterocyclo)alkyl; and

R² is aryl or heterocyclo.

The compounds of formula I are useful in the treatment of
arrhythmia. The invention is also concerned with pharmaceutical
compositions comprising one or more of the novel compounds as an
20 active antiarrhythmic agent either alone or in combination with other
cardiovascular agents such as a β -blocker or other antiarrhythmic
agent; and a method of treating arrhythmia by administration of one of
the novel compounds or compositions thereof to a patient in need of such
treatment.

25

Detailed Description of the Invention

Definition of Terms

Listed below are definitions of various terms used to describe the
30 compounds of the instant invention. These definitions apply to the terms

as they are used throughout the specification (unless they are otherwise limited in specific instances) either individually or as part of a larger group.

5 The term "alkyl" refers to both straight and branched chain groups having 1 to 8 carbon atoms, preferably 1 to 5 carbons, such as methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, the various branched chain isomers thereof, such as isopropyl, t-butyl, isobutyl, isohexyl, 4,4-dimethylpentyl, 2,2,4-trimethylpentyl and the like; as well as such groups substituted by, one or more substituents such as halo,
10 alkoxy, amino, substituted amino, aryl, cycloalkyl, hydroxy, alkanoylamino, arylcarbonylamino, nitro, cyano, thiol, alkylthio and the like.

The term "alkoxy" refers to alkyl-O-.

The term "alkylthio" refers to alkyl-S-.

15 The term "alkenyl" refers to any of the above alkyl groups further containing at least one carbon to carbon double bond.

The term "alkynyl" refers to any of the above alkyl groups further containing at least one carbon to carbon triple bond.

The term "alkanoyl" refers to alkyl-C(O)-

20 The term "cycloalkyl" refers to saturated cyclic hydrocarbon groups containing 3 to 8 ring carbons optionally substituted with one or more substituents such as alkyl or hydroxy.

The term "halogen" or "halo" refers to chlorine, bromine, iodine and fluorine.

25 The term "aryl" refers to monocyclic or bicyclic aromatic hydrocarbon groups having 6 to 12 carbon atoms in the ring portion, such as phenyl, 1-naphthyl, 2-naphthyl, phenanthrene or dihydrophenanthrene; or such groups substituted with one or more substituents such as alkyl, alkenyl, alkynyl, alkylthio, alkoxy, halo,
30 nitro, cyano, hydroxy, amino, substituted amino, phenyl, -C(O)-phenyl, substituted phenyl, -C(O)-substituted amino, heterocycle, carboxylic acid or carboxylic ester.

The term "aryl" also includes those groups listed above fused to a five- or six-membered ring which optionally contains an oxygen, sulfur or nitrogen atom. The five- or six-membered ring may further optionally be substituted with for example, alkyl or -phenyl-CF₃.

5 The term "heterocyclo" or "hetero" refers to fully saturated or unsaturated rings of five or six atoms containing one or two oxygen and/or sulfur atoms and/or one to four nitrogen atoms provided that the total number of hetero atoms in the ring is four or less. Exemplary monocyclic heterocyclo groups include 2- and 3-thienyl, 2- and 3-furyl, 2-,
10 3- and 4-pyridyl and imidazolyl.

The term heterocyclo or hetero also includes bicyclic rings wherein the five- or six-membered ring containing oxygen and/or sulfur and/or nitrogen atoms as defined above is fused to a benzene ring and the bicyclic ring is attached by way of an available atom.

15 Exemplary bicyclic hetero groups include 4-, 5-, 6- or 7-indolyl, 4-, 5-, 6- or 7-isoindolyl, 5-, 6-, 7- or 8-quinolinyl, 5-, 6-, 7- or 8-isoquinolinyl, 4-, 5-, 6- or 7-benzothiazolyl, 4-, 5-, 6- or 7-benzoxazolyl, 4-, 5-, 6- or 7-benzimidazolyl, 4-, 5-, 6- or 7-benzoxadiazolyl and 4-, 5-, 6- or 7-benzofuranyl.

20 The term heterocyclo or hetero also includes such monocyclic and bicyclic rings wherein an available atom is substituted by one or more substituents such as alkyl, aryl, alkylthio, alkoxy, halo, nitro, keto, cyano, hydroxy, azo, oxo, thiazo, amino, substituted amino, carboxylic acid, carboxylic ester, or alkoxy further substituted with a carboxylic
25 acid or a five- to eight-membered ring optionally containing 1 to 4 heteroatoms selected from oxygen, nitrogen and sulfur, optionally substituted by groups such as alkyl or halogen.

The term "substituted amino" refers to a group of the formula -NZ²Z³ wherein Z² is hydrogen, alkyl, cycloalkyl, aryl, morpholinylalkyl,
30 heterocyclo or (heterocyclo)alkyl and Z³ is hydrogen, alkyl, cycloalkyl or aryl further substituted with a carboxylic acid or carboxylic ester, provided that when Z² is hydrogen, then Z³ is other than hydrogen; or Z²

and Z³ taken together with the nitrogen atom to which they are attached are 1-pyrrolidinyl, 1-piperidinyl, 1-azepinyl, 4-morpholinyl, 4-thiamorpholinyl, 1-piperazinyl, 4-alkyl-1-piperazinyl, 4-arylalkyl-1-piperazinyl, 4-diarylalkyl-1-piperazinyl; or 1-pyrrolidinyl, 1-piperidinyl, or 1-azepinyl, optionally substituted with alkyl, alkoxy, alkylthio, halo, aryl or hydroxy.

Throughout the specification, groups and substituents thereof are chosen to provide stable moieties and compounds.

The compounds of formula I may form salts which are also within the scope of this invention. Pharmaceutically acceptable (i.e., non-toxic, physiologically acceptable) salts are preferred, although other salts are also useful, e.g., in isolating or purifying the compounds of this invention.

The compounds of formula I may form salts with alkali metals such as sodium, potassium and lithium, with alkaline earth metals such as calcium and magnesium, with organic bases such as dicyclohexylamine, tributylamine, pyridine and amino acids such as arginine, lysine and the like. Such salts may be obtained, for example, by exchanging the carboxylic acid protons, if they contain a carboxylic acid, in compound I with the desired ion in a medium in which the salt precipitates or in an aqueous medium followed by evaporation. Other salts can be formed as known to those having ordinary skill in the art.

The compounds of formula I may form salts with a variety of organic and inorganic acids. Such salts include those formed with hydrogen chloride, hydrogen bromide, methanesulfonic acid, sulfuric acid, acetic acid, trifluoroacetic acid, maleic acid, benzenesulfonic acid, toluenesulfonic acid and various others (e.g., nitrates, phosphates, borates, tartrates, citrates, succinates, benzoates, ascorbates, salicylates and the like). Such salts may be formed by reacting compound I in an equivalent amount of the acid in a medium in which the salt precipitates or in an aqueous medium followed by evaporation.

In addition, zwitterions ("inner salts") may be formed.

A compound of the formula I may also have prodrug forms. Any compound that will be converted *in vivo* to provide the bioactive agent (i.e., the compound of formula I) is a prodrug within the scope and spirit of the invention.

5 Various forms of prodrugs are well known in the art. For examples of such prodrug derivatives, see:

a) *Design of Prodrugs*, edited by H. Bundgaard, (Elsevier, 1985);

b) *Methods in Enzymology*, Vol. 42, 309-396, edited by K. Widder et al. (Academic Press, 1985);

10 c) *A Textbook of Drug Design and Development*, edited by Krogsgaard-Larsen and H. Bundgaard, Chapter 5, "Design and Application of Prodrugs," by H. Bundgaard, 113-191 (1991);

d) *Advanced Drug Delivery Reviews*, H. Bundgaard, 8, 1-38 (1992);

e) *Journal of Pharmaceutical Sciences*, H. Bundgaard et al., 77, 285
15 (1988); and

f) *Chem Pharm Bull*, N. Kakeya et al., 32, 692 (1984).

It should further be understood that solvates (e.g., hydrates) of the compounds of formula I are also within the scope of the present invention. Methods of solvation are generally known in the art.

20 All stereoisomers of the compounds of the instant invention are contemplated, either in admixture or in pure or substantially pure form. The compounds of the present invention can have asymmetric centers at any of the carbon atoms including any one of the R substituents. Consequently, compounds of formula I can exist in diastereomeric
25 forms or in mixtures thereof. The below described processes can utilize racemates, enantiomers or diastereomers as starting materials. When diastereomeric products are prepared, they can be separated by conventional methods for example, chromatography or fractional crystallization.

30

Use and Utility

The compounds of formula I are useful in the treatment of arrhythmia. More specifically, the compounds of the present invention have the pharmacological properties required for the antiarrhythmic agents of Class III.

Class III agents increase myocardial refractoriness via a prolongation of cardiac action potential duration. Theoretically, prolongation of the cardiac action potential can be achieved by enhancing inward currents (i.e. Na^+ or Ca^{2+} currents; hereinafter I_{Na} and I_{Ca} respectively) or by reducing outward repolarizing potassium (K^+) currents. The delayed rectifier (I_{K}) K^+ current is the main outward current involved in the overall repolarization process during the action potential plateau, whereas the transient outward (I_{to}) and inward rectifier (I_{K1}) K^+ current are responsible for the rapid initial and terminal phases of repolarization, respectively. Cellular electrophysiologic studies have demonstrated that I_{K} consists of two pharmacologically and kinetically distinct K^+ current subtypes, I_{Kr} (rapidly activating and deactivating) and I_{Ks} (slowly activating and deactivating).

Most Class III agents that are known to be in development predominantly block I_{Kr} . These agents have a potential liability in that they have an enhanced risk of proarrhythmia at slow heart rates. The compounds of the present invention prolong the myocardial action potential in vitro without a significant depression of the V_{max} and with the prolongation of Qtc-interval in anesthetized dogs. In addition the compounds of the present invention selectively block I_{Ks} . The preferred compounds of the present invention are those which have selectivity of $I_{\text{Ks}}:I_{\text{Kr}}$ greater than or equal to 5.

The compounds of the present invention are effective in treating and preventing all types of arrhythmias including ventricular and atrial (supraventricular) arrhythmias. The compounds of the present invention are especially useful to control reentrant arrhythmias and prevent sudden death due to the ventricular fibrillation.

In the novel method of this invention of treating arrhythmia, a novel compound or pharmaceutically acceptable salt thereof, is administered in an amount ranging from about 0.0001 to about 20 mg. per kg of body weight per day, preferably from about .001 to about 10 mg per kg of body weight per day in a single dose or in 2 to 4 divided doses.

The novel compounds of this invention can be administered as the sole active ingredient or in combination with other antiarrhythmic agents or other cardiovascular agents.

The compounds, or pharmaceutically acceptable salts thereof, of the present invention, in the described dosages, are administered orally, intraperitoneally, subcutaneously, intramuscularly, transdermally, sublingually or intravenously. They are preferably administered orally, for example in the form of tablets, troches, capsules, elixirs, suspensions, syrups, wafers, chewing gum, or the like prepared by art recognized procedures. The amount of active compound in such therapeutically useful compositions or preparations is such that a suitable dosage will be obtained.

Preferred Moieties

The preferred compounds of the present invention are those compounds of formula I where:

X is oxygen or N-CN;

Y is a single bond or -C=C-; and

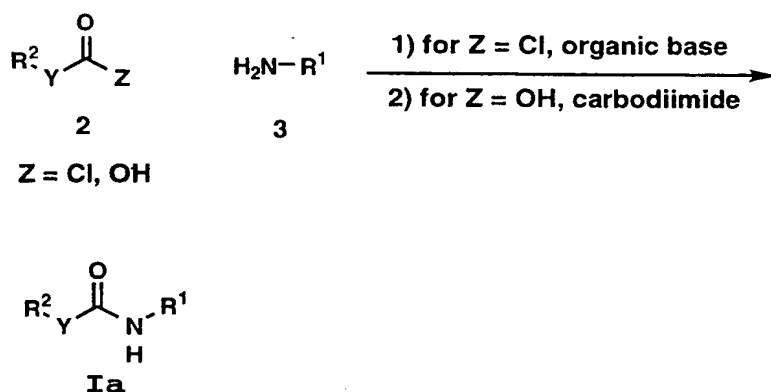
R¹ is alkyl, cycloalkyl, (aryl)alkyl, (cycloalkyl)alkyl, or (substituted amino)alkyl.

Process of Preparation

The compounds of the instant invention may be obtained by methods exemplified by the following descriptions.

The compounds of formula I wherein X is O and Y is a single bond or -C=C- (compounds of formula Ia), can be prepared as outlined in Scheme 1 below.

Scheme 1



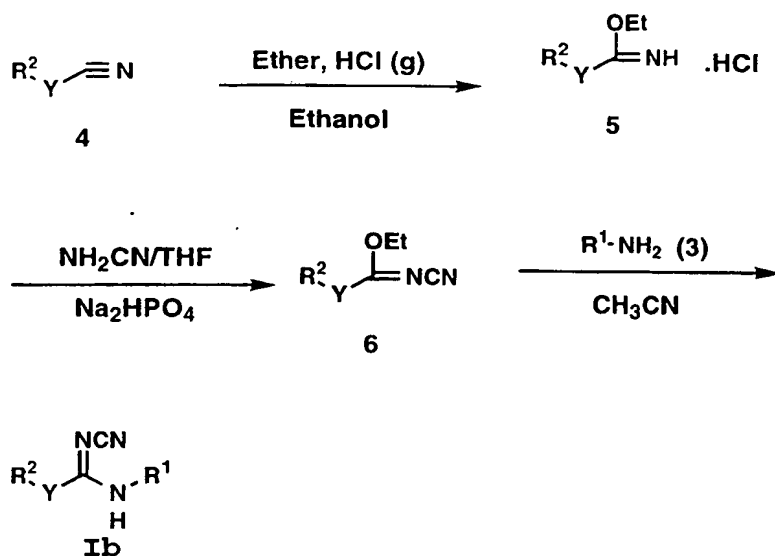
The acid chloride (Z = Cl) of formula 2 is reacted with an amine of formula 3 in the presence of a base such as triethylamine and an organic solvent such as dichloromethane, tetrahydrofuran or dimethylformamide etc. to form the compounds of formula Ia.

Alternatively, compounds of formula Ia can be prepared from an acid of formula 2 (Z = OH) and an amine of formula 3 in an organic solvent in the presence of a carbodiimide such as dicyclohexylcarbodiimide or 1-(3-dimethylaminopropyl)-2-ethylcarbodiimide hydrochloride (WSC).

Compounds of formula I wherein X is S, can be prepared from compounds of formula I wherein X is O by treatment with a thionating agent such as phosphorus pentasulfide (P₄S₁₀) or the Lawesson's reagent. Compounds of formula 2 and 3 are commercially available or they can be prepared by methods described in the literature.

The compounds of formula I wherein X is NCN and Y is a single bond or -C=C- (i.e., compounds of formula Ib), can be prepared as outlined in Scheme 2.

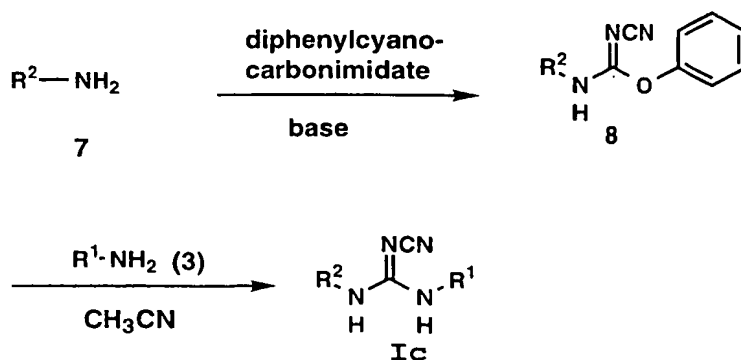
Scheme 2



The nitrile of formula 4 is converted to the imino ether of formula 5 by treatment with hydrochloric acid in ethanol. The imino ether 5 is then reacted with cyanamide to provide a compound of formula 6 which on reaction with an amine of formula 3 provides the compounds of formula Ib. Compounds of formula 4 are commercially available or they can be prepared by methods described in the literature.

The compounds of formula I wherein X is NCN and Y is NH (i.e., compounds of formula Ic), can be prepared as outlined in Scheme 3.

Scheme 3

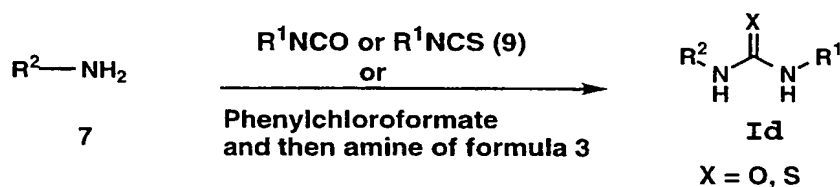


The amine of formula 7 is converted to the phenyl ester of formula 8 by treatment with diphenylcyanocarbonimidate in the presence of a base (e.g., sodium hydride, diisopropylethyl amine). The compound of formula 8 is converted to the compounds of formula Ic by treatment with an amine of formula 3 in an organic solvent. Compounds of formula 7 are commercially available or they can be prepared by methods described in the literature.

Alternatively, compounds of formula I wherein X is NCN and Y is NH, can be prepared by methods similar to those described in the literature such as that by Atwal et. al. Tetrahedron Letters, Vol. 30, pp 7313-7316 (1989) and references therein.

The compounds of formula I wherein X is O or S and Y is NH (i.e., compounds of formula Id), can be prepared as outlined in Scheme 4.

15

Scheme 4

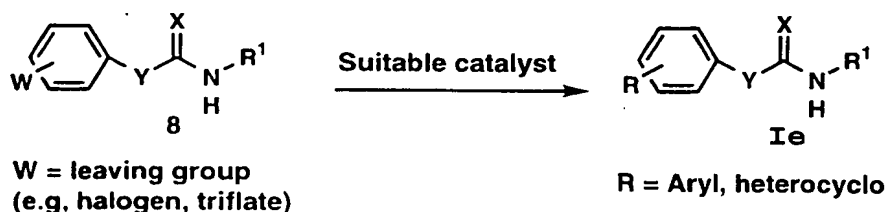
The amine of formula 7 is reacted with an isocyanate or isothiocyanate of formula 9 in an organic solvent to provide the compounds of formula Id.

Alternatively, the compounds of formula I wherein X is O or S and Y is NH, can be prepared by sequential treatment of an amine of formula 7 with an arylchloroformate (e.g., phenylchloroformate) and an amine of formula 3.

Compounds of formula I wherein X is S, can be prepared from compounds of formula I wherein X is O by treatment with a standard thionating agent such as phosphorus pentasulfide (P₄S₁₀) or the Lawesson's reagent. Compounds of formula 9 are commercially available or they can be prepared by conventional methods described in the literature.

The compounds of formula I wherein R² contains an aryl or a heterocyclo substituent (formula Ie) may be prepared according to Scheme 5.

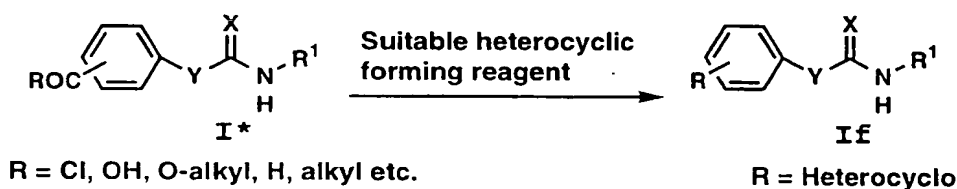
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Scheme 5

Compounds of formula Ie can be prepared from compounds of formula 8 wherein W is a suitable leaving group such as halogen or triflate by treatment with a heterocyclo group or an aryl ring containing a suitable reacting group such as an amino group, boronic acid and trialkyltin in the presence of a suitable catalyst (base, palladium etc.). Compounds of formula 8 are prepared by methods described in Schemes 1 to 4.

The compounds of formula I wherein R² contains a heterocyclic substituent (oxazole, imidazole, oxadiazole, thiadiazole etc.), can also be prepared from compounds of formula I* wherein R² contains a suitable heterocyclo forming group such as an acid or a derivative thereof as described in Scheme 6.

20

Scheme 6

Compounds of formula If are prepared by treatment with a suitable heterocyclo forming reagent (amidine, hydroxyamidine etc.). This

transformation can also be accomplished in a stepwise fashion by modification of methods described in the literature.

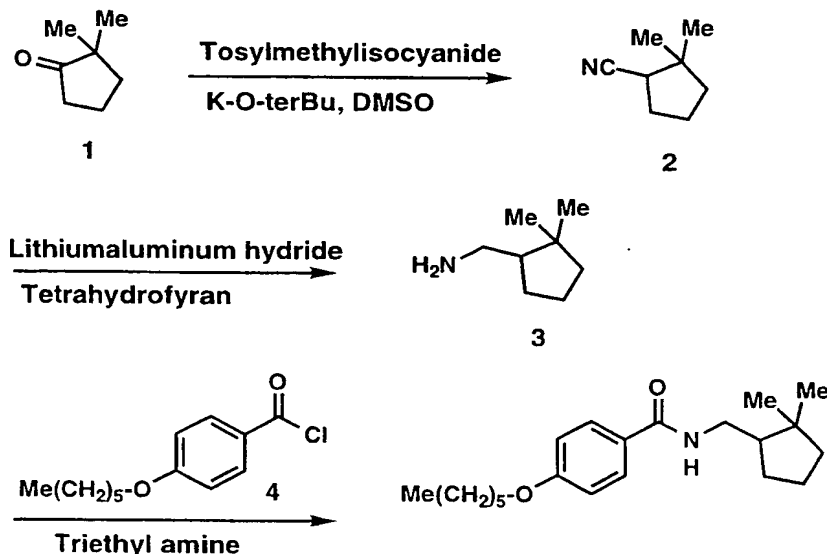
All other compounds of formula I may be prepared by modification of the procedures discussed herein as known by those having ordinary skill in the art. The intermediates used to prepare compounds of formula I are described herein or may be derived from known compounds by those having ordinary skill in the art or are commercially available.

10 Examples

The following examples and preparations describe the manner and process of making and using the invention and are illustrative rather than limiting. It should be understood that there may be other embodiments which fall within the spirit and scope of the invention as defined by the claims appended hereto.

Example 1

(+)-N-[(2,2-Dimethylcyclopentyl)methyl]-4-hexyloxybenzamide and (-)-N-[(2,2-Dimethylcyclopentyl)methyl]-4-hexyloxybenzamide



A. 2,2-Dimethylcyclopentanecarbonitrile

Tosylmethyl isocyanide (3.76 g, 19.3 mmol) was dissolved in dimethylsulfoxide (15 mL) and cooled to 0°C. Solid potassium *t*-butoxide (6.3 g, 59.2 mmol) was added and a thick brown precipitate formed. The reaction mixture was warmed to room temperature and 2,2-dimethylcyclopentanone (2.0 mL, 16.0 mmol) was added in anhydrous methanol (680 µL) and the reaction mixture was stirred for 30 hours. It was diluted with ethyl acetate, acidified with hydrochloric acid and extracted with hexanes. The extracts were dried over magnesium sulfate, filtered and the solvent was removed to provide a brown oil which was purified by chromatography on silica gel eluted with 0-50% dichloromethane in hexanes to yield a yellow oil (612 mg, 31%).

B. 2,2-Dimethylcyclopentanemethanamine

Compound 2 (600 mg, 4.87 mmol) was dissolved in tetrahydrofuran (10 mL) and cooled to 0°C. Lithium aluminum hydride (185 mg, 4.87 mmol) was added and the mixture was stirred for 17 hours. The reaction mixture was then quenched by addition of sodium sulfate decahydrate, filtered through celite and the celite washed with ether. To the solution was added hydrogen chloride (3 mL, 3.9 M in dioxane) and the solvent was removed. The residue was dissolved in water and lyophilized to yield a white solid (613 mg, 77%).

C. (+)-N-[(2,2-Dimethylcyclopentyl)methyl]-4-hexyloxybenzamide and (-)-N-[(2,2-Dimethyl-cyclopentyl)methyl]-4-hexyloxybenzamide:

To compound 3 (300 mg, 1.83 mmol) in dichloromethane (8 mL) was added triethylamine (766 µL, 5.50 mmol) and 4-hexyloxybenzoyl chloride (4) (430 µL, 1.92 mmol) and the reaction mixture was stirred for 24 hours at room temperature. The reaction mixture was diluted with ethyl acetate and washed with hydrochloric acid, saturated sodium bicarbonate and brine. The organic layer was dried over magnesium

sulfate, filtered and the solvent was removed to yield a white solid (600 mg). Purification by preparative HPLC (Chiracel OD, 50 x 500 mm; 10% isopropanol, hexane; 90 mL/min) provided 99 mg of a faster moving isomer, 342 mg of mixed fraction and 65 mg of a slower moving isomer.

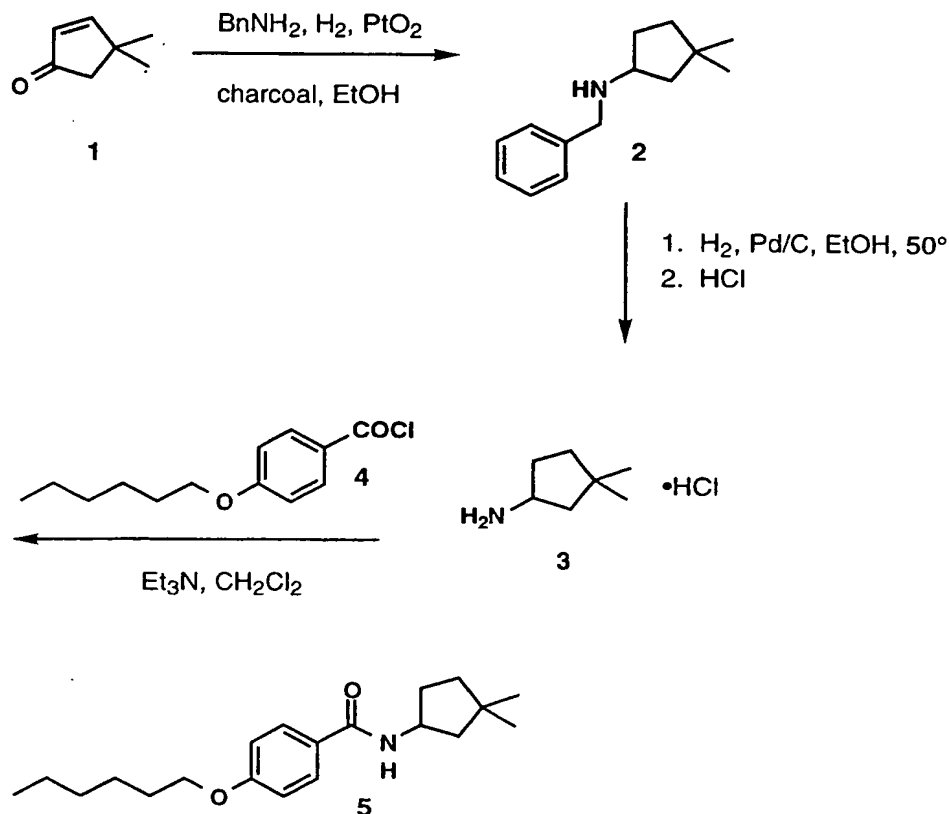
- 5 Further purification of the faster moving isomer by flash chromatography on silica gel eluted with 10% acetone in hexanes provided (+)-N-[(2,2-dimethylcyclopentyl)methyl]-4-hexyloxybenzamide as a white solid (90 mg): mp 68-69°C; $[\alpha]_D$ (MeOH, c 0.55) +20.7°.

Analysis calculated for $C_{21}H_{33}NO_2$: C, 76.09; H, 10.03; N, 4.24. Found:

- 10 C, 75.99; H, 10.11; N, 4.11.

Further purification of the slower moving isomer by flash chromatography on silica gel eluting with 10% acetone in hexane provided (-)-N-[(2,2-dimethylcyclopentyl)-methyl]-4-hexyloxybenzamide as a white solid (57 mg): mp 69-70°C; $[\alpha]_D$ (MeOH, c 0.55) -18.5°;

- 15 Analysis calculated for $C_{21}H_{33}NO_2$: C, 76.09; H, 10.03; N, 4.24. Found: C, 75.95; H, 10.12; N, 4.13.

Example 1a**N-(3,3-Dimethylcyclopentyl)-4-hexyloxybenzamide****A. N-(3,3-Dimethylcyclopentyl)benzenemethanamine**

4,4-Dimethyl-2-cyclopenten-1-one (804 mg, 7.30 mmol) and benzylamine (877 μL , 8.03 mmol) were dissolved in 15 mL of ethanol. Charcoal (400 mg) and platinum oxide (Adams' catalyst, 331 mg, 1.46 mmol) were added and the mixture was stirred under hydrogen (balloon) for 24 hours. The reaction was purged with argon and diluted with ethyl acetate. The mixture was loaded onto silica gel and purified by flash chromatography on silica gel eluted with 5% triethylamine and hexane to provide 1.24 g (84%) of a clear colorless oil.

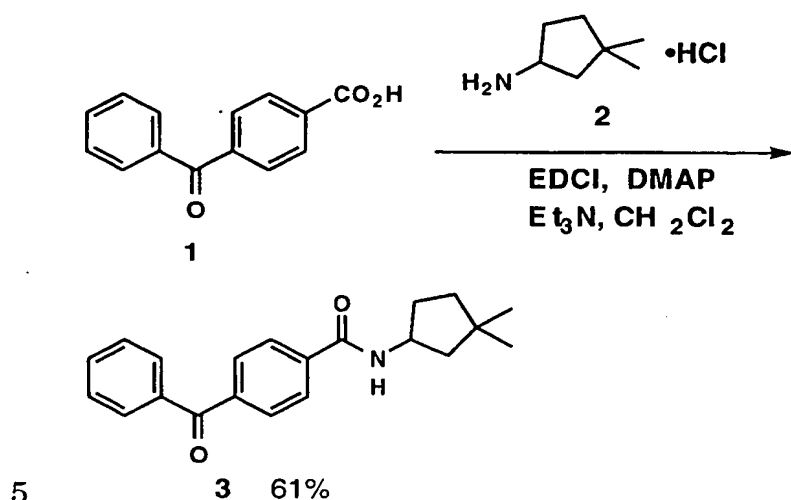
B. 3,3-Dimethylcyclopentanamine hydrochloride

The amine (2, 130 mg, 0.64 mmol) was dissolved in 2 mL of ethanol. Palladium (5% on charcoal, 13 mg) was added and the reaction was stirred under hydrogen (balloon) at 50°C for 10 hours then at room temperature for 10 hours. The mixture was diluted with ether then filtered through Celite. Hydrogen chloride (3.86 M, 300 µL) was added and the solvent removed to yield 87 mg (91%) of a white solid.

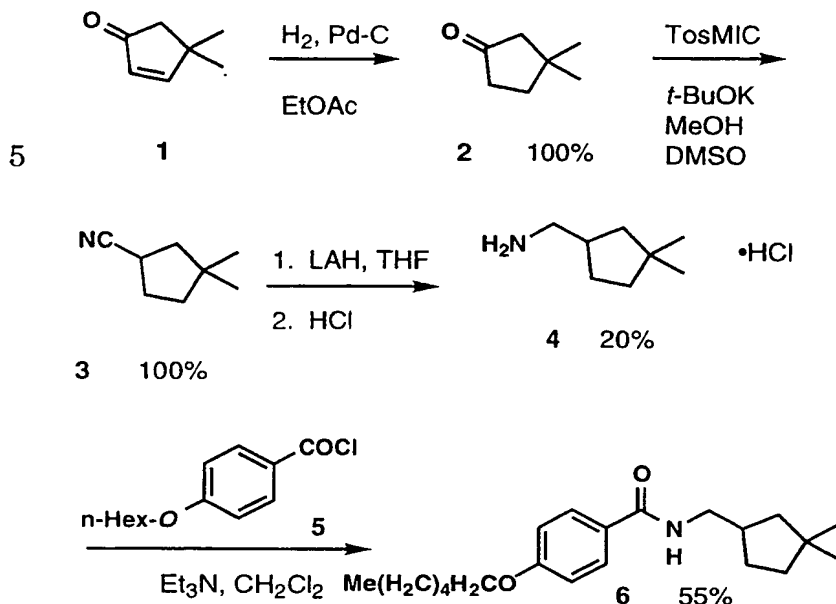
C. N-(3,3-Dimethylcyclopentyl)-4-hexyloxybenzamide

The amine hydrochloride (3, 87 mg, 0.58 mmol) was dissolved in 2 mL dichloromethane. Triethylamine (244 µL, 1.75 mmol) and 4-hexyloxybenzoyl chloride (155 µL, 0.70 mmol) were added and the mixture was stirred for 4 hours at room temperature. The reaction was diluted with ethyl acetate and washed with hydrochloric acid (1.0 M, aq.), sodium bicarbonate (sat'd., aq.) and sodium chloride (sat'd., aq.). The organic layer was dried over magnesium sulfate, filtered and the solvent removed. Purification by flash chromatography on silica gel eluted with 10% acetone, hexane provided 138 mg (75%) of a white solid. mp 82-83°C; Anal calc'd for C₂₀H₃₁NO₂: C, 75.67; H, 9.84; N, 4.41.

Found: C, 75.86; H, 10.10; N, 4.37.

Example 1c**N-(3,3-Dimethylcyclopentyl)-4-benzoylbenzamide**

4-Benzoylbenzoic acid (0.20 mmol) and the title B compound of example 1b (0.22 mmol) were dissolved in 1 mL of THF and triethylamine (0.22 mmol) and 4-dimethylamino-pyridine (0.02 mmol) were added. 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (1.6 ML, 0.15 M in dichloromethane, 0.24 mmol) was added and the mixture was allowed to stand for 48 hours. Purification by preparative HPLC (YMC ODS-A S% 30 x 250 mm column, 50-90% methanol water with 0.10% TFA gradient over 20 minutes, 25 mL/minute flow) provided 40 mg (61%) of a white solid. C₂₁H₂₃NO₂: m/z = 322 (M+H).

Example 1d**N-[(3,3-Dimethylcyclopentyl)methyl]-4-(hexyloxy)benzamide****A. 3,3-Dimethylcyclopentanone**

4,4-Dimethyl-2-cyclopenten-1-one (**1**, 1.0 g, 9.08 mmol) was dissolved in 5 mL of ethyl acetate and palladium (10% on carbon, 50 mg) was added. The mixture was stirred under a hydrogen atmosphere (balloon) for 23 hours. The reaction was then filtered through Celite and the Celite washed with ether. The solvent was removed by distillation at atmospheric pressure to provide 1.11 g (>100%) of a clear colorless oil that was used without further purification.

B. 3,3-Dimethylcyclopentanecarbonitrile

Solid potassium *t*-butoxide (3.57 g, 31.8 mmol) was dissolved in 10 mL of anhydrous dimethylsulfoxide. Tosylmethyl isocyanide (2.13 g, 10.9 mmol) was dissolved in 8 mL of dimethylsulfoxide and added to the potassium-*t*-butoxide. A thick brown precipitate formed. The ketone (**2**, assumed 9.08 mmol) was added in 386 μ L of anhydrous methanol. The

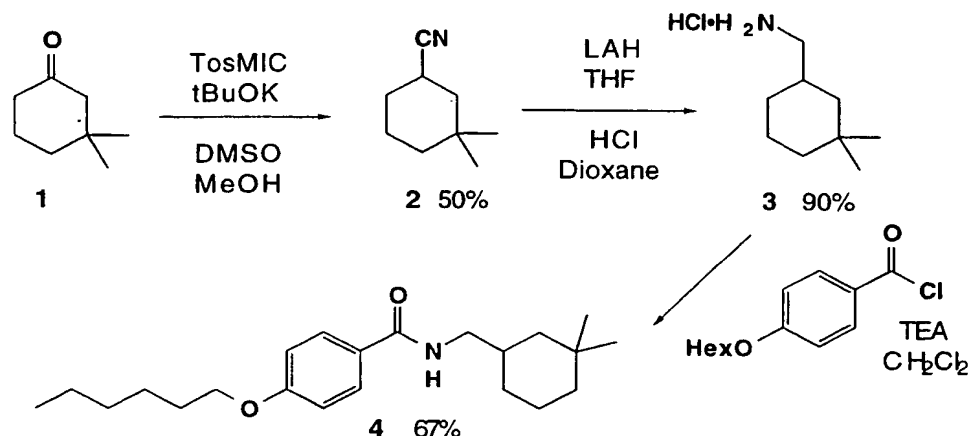
mixture was heated to 45°C and stirred for 24 hours. The reaction was diluted with water (100 mL), acidified to pH 3, and extracted with pentane. The extracts were dried over magnesium sulfate, filtered and the solvent removed to provide a yellow oil (>100%) that was used without purification.

C. 3,3-Dimethylcyclopentanemethanamine hydrochloride

The nitrile (3, assumed 9.08 mmol) was dissolved in 10 mL of anhydrous tetrahydrofuran and cooled to 0°C. Lithium aluminum hydride (335 mg, 9.08 mmol) was added and the mixture was stirred for 20 hours. The reaction was then quenched by addition of sodium sulfate decahydrate, filtered through celite and the celite washed with ether. Hydrogen chloride (2.5 mL, 4 M in dioxane) was added and the solvent removed to provide a brown oil. This oil was dissolved in water, filtered and lyophilized to yield 296 mg (20%) of a brown solid.

D. N-[(3,3-Dimethylcyclopentyl)methyl]-4-(hexyloxy)benzamide

The amine hydrochloride (4, 100 mg, 0.61 mmol) was dissolved in 2 mL of dichloromethane. Triethylamine (255 µL, 1.83 mmol) and 4-hexyloxybenzoyl chloride (5, 136 µL, 0.61 mmol) were added and the mixture was stirred for 6 hours at room temperature. The reaction was diluted with ethyl acetate and washed with hydrochloric acid (1.0 M, aq.), sodium bicarbonate (sat'd., aq.) and sodium chloride (sat'd., aq.). The organic layer was dried over magnesium sulfate, filtered and the solvent removed to provide 210 mg of a brown solid. Purification by flash chromatography on silica gel eluting with 5% acetone, hexane provided 111 mg (55%) of a white powder. mp 85-86°C; Anal. calc'd for C₂₁H₃₃NO₂: C, 76.09; H, 10.03; N, 4.23. Found: C, 76.02; H, 10.23; N, 4.20.

Example 1e**N-[(3,3-Dimethylcyclohexyl)methyl]-4-(hexyloxy)benzamide**

5

A. 3,3-Dimethylcyclohexanecarbonitrile

Tosylmethyl isocyanide (3.0 g, 15.4 mmol) was dissolved in 15 mL of anhydrous dimethylsulfoxide and the solution was cooled to 5 - 10°C in an ice water bath. Potassium *tert*-butoxide (5.03g, 44.8 mmol) was then added and the solution turned dark brown in color. The reaction was allowed to warm to room temperature and stirred for 1 hour. A solution of the cyclohexanone (1) (1.62 g, 12.8 mmol) in anhydrous methanol (685 µL) was added to the reaction via syringe and the reaction was stirred at room temperature under argon for 40 hours. The reaction was then diluted with 25 mL of water and placed in an ice water bath and the reaction was made acidic by the dropwise addition of aqueous HCl (6.0 M, aq.). The aqueous phase was extracted 2 times with hexane, and the organic phases were combined and dried over MgSO₄, filtered and concentrated to provide 2.24 g of a yellow mixture of oil and solid. The crude mixture was purified by chromatography on silica gel (eluted with 0-50% dichloromethane in hexanes) and the fractions were concentrated and monitored by IR to find 870 mg (50%) of the nitrile product (2) as a transparent oil.

B. 3,3-Dimethylcyclohexanemethanamine hydrochloride

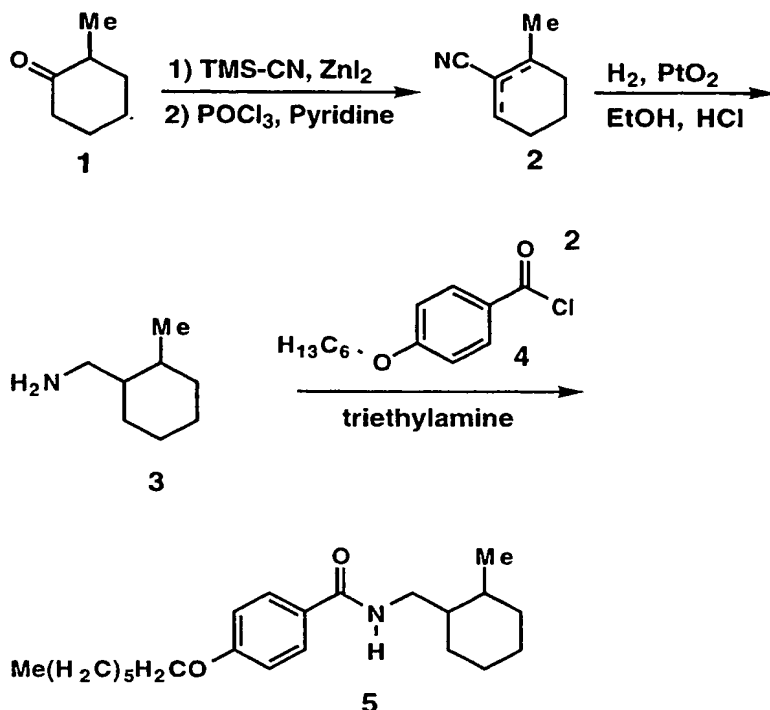
The nitrile (2) (410 mg, 2.99 mmol) was dissolved in 6 mL of anhydrous tetrahydrofuran and the resulting solution was cooled to 0°C on an ice water bath. Solid lithium aluminum hydride (114 mg, 3.0 mmol) was then slowly added to the reaction in small portions. Upon full addition, the bath was removed and the reaction was stirred at room temperature under argon for 16 hours. The reaction was then cooled on an ice water bath and solid Na₂SO₄ • 10 H₂O was slowly added until gas evolution ceased. The reaction was diluted in 10 mL of Et₂O and several drops of water were added to produce a filterable solid. The reaction was then filtered through a pad of Celite and 1.5 mL of 4.0 M HCl in dioxane was added to the filtrate. The cloudy reaction solution was stirred for 15 minutes and then concentrated *in vacuo* to provide 477 mg (90%) of a white solid.

C. N-[(3,3-Dimethylcyclohexyl)methyl]-4-(hexyloxy)-benzamide

The amine (3) was dissolved in 5 mL of anhydrous dichloromethane and triethylamine (171 µL, 1.23 mmol) was added at room temperature under argon. The solution was cooled in an ice water bath and hexyloxybenzoyl chloride (134 µL, 0.61 mmol) was added via syringe and the reaction was allowed to warm to room temperature and stirred for 64 hours. The reaction was partitioned between HCl (1.0 M, aq) and dichloromethane and the organic phase was washed with NaHCO₃ (sat'd, aq.), dried over MgSO₄, filtered and concentrated to provide 224 mg of a white solid. The solid was purified by flash chromatography on silica gel eluted with 15% acetone, hexane to provide 142 mg (67%) of the desired amide. Analysis calculated for C₂₂H₃₅NO₂: C, 76.48; H, 10.21; N, 4.05. Found: C, 76.42; H, 10.27; N, 3.98.

Example 1f

4-(Hexyloxy)-N-[(2-methylcyclohexyl)methyl]benzamide



A. 6-methyl-1-cyclohexenenitrile and 2-methyl-1-cyclohexenenitrile

Trimethylsilyl cyanide (8.0 mL, 60.0 mmol) was added dropwise over a 5 minute period to a suspension of 2-methylcyclopentanone (1) (6.1 mL, 50.0 mmol) and zinc iodide (399 mg, 1.25 mmol) in methylene chloride (25 mL). After stirring for 3 hours, a mixture of phosphorous oxychloride (11.2 mL, 120 mmol) and pyridine (60 mL) was added to the reaction. The reaction was heated at 110°C for 23 hours and then cooled to room temperature. The reaction was carefully poured onto 250 g of crushed ice, acidified with 6 N HCl (30 mL), and extracted with ether (3 x 150 mL). The combined organic layers were washed with water (50 mL), saturated NaHCO₃ (50 mL), dried (MgSO₄) and concentrated by distillation. Vacuum distillation (10 mm Hg) provided 5.56 g of a mixture of 6-methyl-1-cyclohexenenitrile and 2-methyl-1-cyclohexenenitrile (bp 86-91°C).

B. ((2-Methylcyclohexyl)methyl)amine hydrochloride

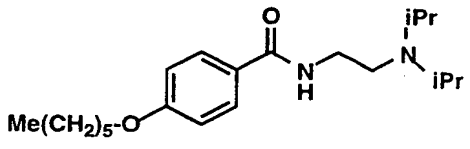
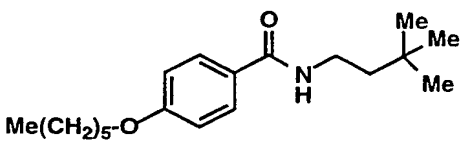
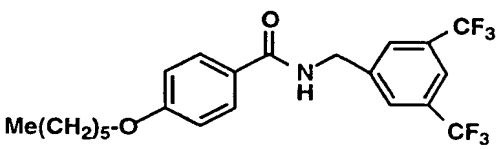
The mixture of title A compounds (2.00 g, 16.5 mmol), PtO_2 (120 mg), and saturated ethanolic HCl (4.5 mL) in ethanol (75 mL) was shaken under a hydrogen atmosphere (55 psi) on a Parr shaker for 3.5 hours. After purging with nitrogen, the reaction was filtered through Celite AFA. The pad was rinsed with ethanol (3 x 15 mL) and the combined filtrates were concentrated *in vacuo*. Trituration of the solid residue with ether (1 x 50 mL, 2 x 20 mL) provided 2.29 g (85%) of ((2-methylcyclohexyl)methyl)amine hydrochloride as a mixture of isomers.

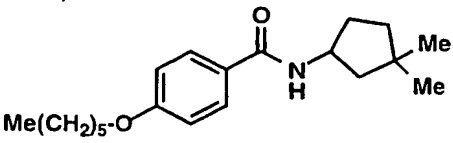
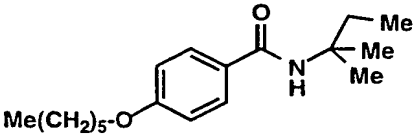
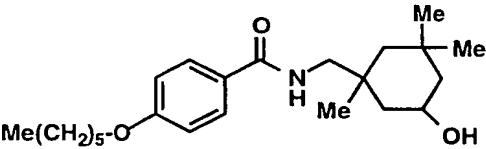
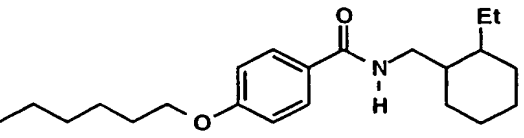
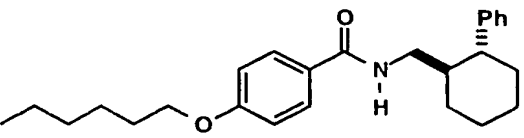
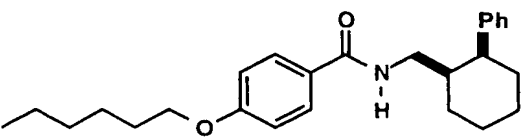
C. 4-(Hexyloxy)-N-[(2-methylcyclohexyl)methyl]benzamide

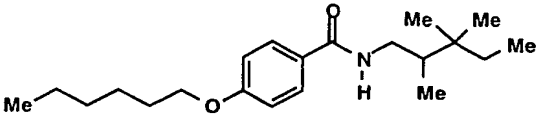
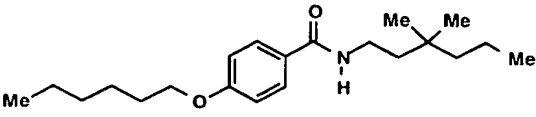
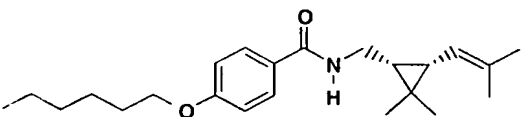
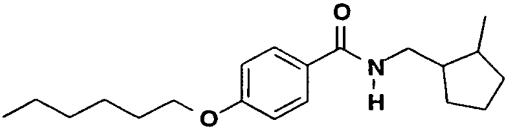
To a suspension of title B compound (213 mg, 1.30 mmol) in THF (6 mL) was added triethylamine 0.42 mL, 3.0 mmol). After stirring for 5 minutes, 4-hexyloxybenzoyl chloride (0.223 mL, 1.00 mmol) was added. After stirring for 4 hours, the reaction was filtered through a Celite pad. The pad was rinsed with ether (4 x 5 mL) and the combined filtrates were concentrated *in vacuo*. Purification by RP-HPLC (YMC SH-365-10 120A S-110 30 x 500 mm column, 40 mL/minute, 90:10 methanol/water (containing 0.1% TFA)) afforded the title compound as a 1:1 mixture of cis- and trans- isomers (43 mg). MS: (ESI) (m/z) 332.

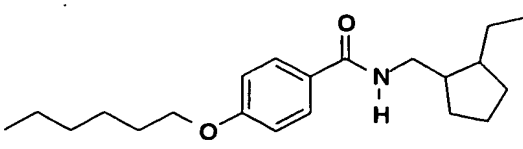
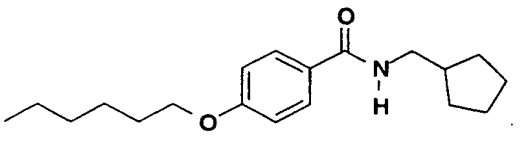
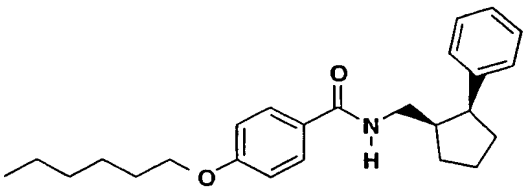
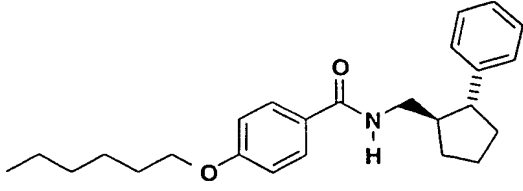
Using the procedure described for the synthesis of the title B compound of Example 1f, ((2-ethylcyclohexyl)-methyl)amine hydrochloride (from 2-ethylcyclohexanone, 71% yield, isomer mixture), ((2-phenylcyclohexyl)methyl)amine hydrochloride (from 2-phenylcyclohexanone, 44% yield, isomer mixture), ((2-methylcyclopentyl)methyl)amine hydrochloride (from 2-methylcyclopentanone, 49% yield, isomer mixture), and ((2-ethylcyclopentyl)methyl)amine hydrochloride (from 2-ethylcyclopentanone, 42% yield, isomer mixture) were prepared.

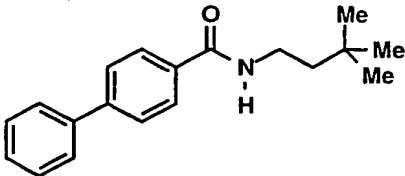
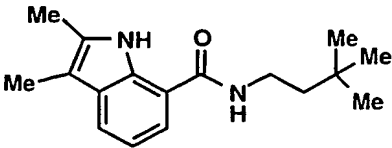
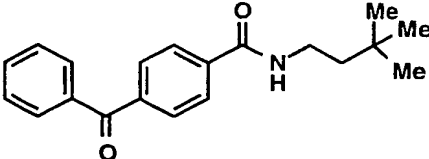
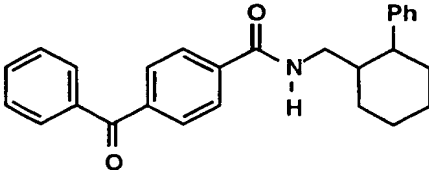
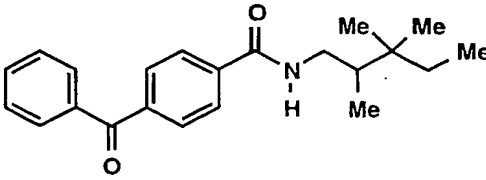
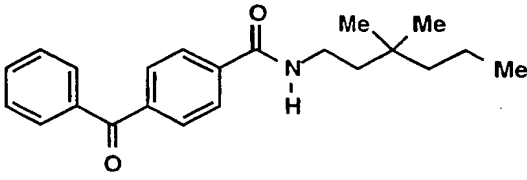
Using methodology analogous to that described for the title compounds of Example 1, the compounds of Examples 2 to 20 were prepared.

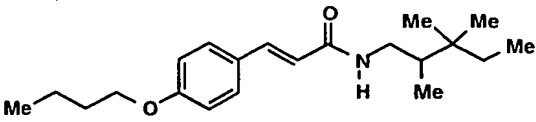
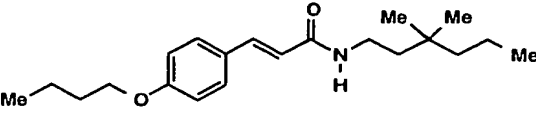
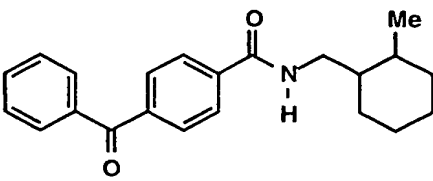
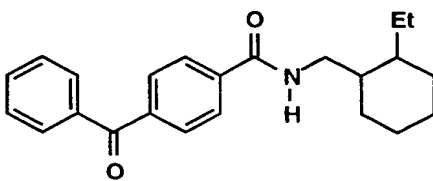
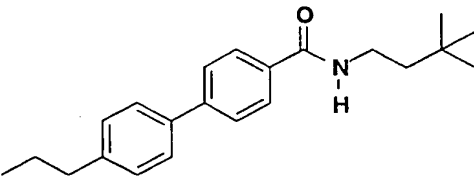
Example #	Structure	Characterization
2		$C_{21}H_{36}N_2O_2 \cdot HCl$: mp 131-134°C. Analysis calculated: C, 65.5; H, 9.69; N, 7.27; Cl, 9.21. Found: C, 65.6; H, 9.79; N, 7.17; Cl, 8.97.
3		$C_{19}H_{31}NO_2$: mp 74-76°C. Analysis calculated: C, 74.71; H, 10.23; N, 4.59. Found: C, 75.01; H, 10.40; N, 4.52.
4		$C_{22}H_{23}F_6NO_2$: mp 105-106°C. Analysis calculated: C, 59.06; H, 5.18; N, 3.13; F, 25.48. Found: C, 58.88; H, 5.15; N, 2.98; F, 25.36.

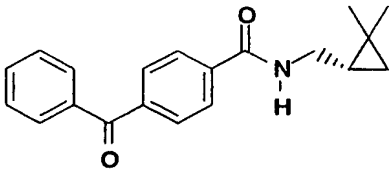
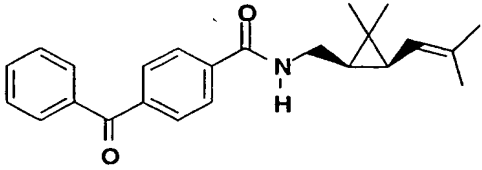
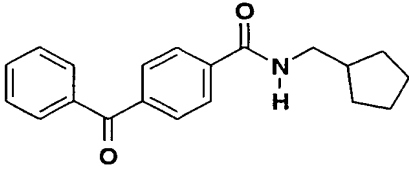
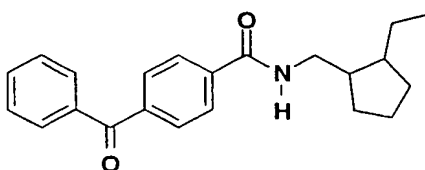
Example #	Structure	Characterization
5		$C_{20}H_{31}NO_2$: mp 82-83°C. Analysis calculated: C, 75.67; H, 9.84; N, 4.41. Found: C, 75.86; H, 10.10; N, 4.37.
6		$C_{18}H_{29}NO_2$: m/e = 291.
7		$C_{23}H_{37}NO_3$: m/e = 375.
7a		$C_{22}H_{35}NO_2$ ESI (m/z) 346 Rf (silica, 25% EtOAc/hex) 0.34.
7b		$C_{26}H_{35}NO_2$ ESI (m/z) 394 Rf (silica, 25% EtOAc/hex) 0.27.
7c		$C_{26}H_{35}NO_2$ ESI (m/z) 394 Rf (silica, 25% EtOAc/hex) 0.28.

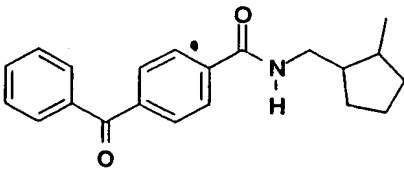
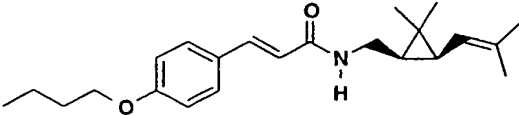
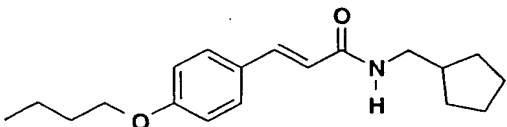
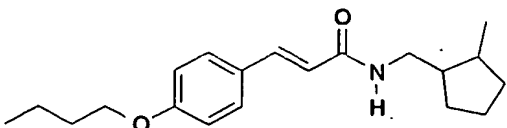
Example #	Structure	Characterization
7d		$C_{21}H_{35}NO_2$ ESI (m/z) 334 Rf (silica, 25% EtOAc/hex) 0.39.
7e		$C_{21}H_{35}NO_2$ ESI (m/z) 334 Rf (silica, 25% EtOAc/hex) 0.41.
7f		$C_{23}H_{35}NO_2$: ESI (m/z) 358 mp 68.0-69.0°C. Analysis calculated: C, 77.27; H, 9.87; N, 3.92. Found: C, 77.02; H, 10.12; N, 3.74.
7g	 60:40 isomer mixture	$C_{20}H_{31}NO_2$: mp 98.0-100.0°C. Analysis calculated: C, 75.67; H, 9.84; N, 4.41. Found: C, 75.57; H, 10.02; N, 4.26.

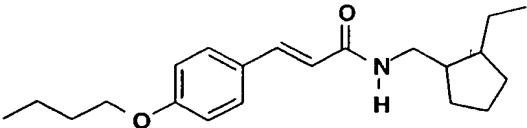
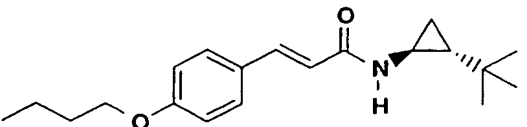
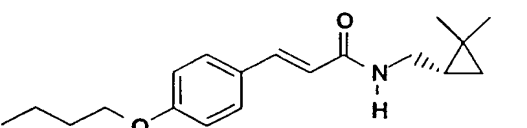
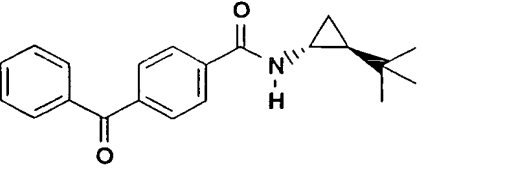
Example #	Structure	Characterization
7h	 <p>50:50 trans:cis mixture</p>	<p>$C_{21}H_{33}NO_2 \cdot 0.14$ H_2O: mp 93.0-95.5°C. Analysis calculated: C, 75.50; H, 10.04; N, 4.19. Found: C, 75.50; H, 10.20; N, 3.94.</p>
7i		<p>$C_{19}H_{29}NO_2 \cdot 0.17$ H_2O: mp 107.0-109.5°C. Analysis calculated: C, 74.45; H, 9.65; N, 4.57. Found: C, 74.45; H, 9.89; N, 4.71.</p>
7j		<p>$C_{25}H_{33}NO_2$: mp 87.0-88.0°C. Analysis calculated: C, 79.12; H, 8.76; N, 3.69. Found: C, 79.09; H, 8.67; N, 3.58.</p>
7k		<p>mp 77.0-78.0°C ESI (m/z) 380.</p>

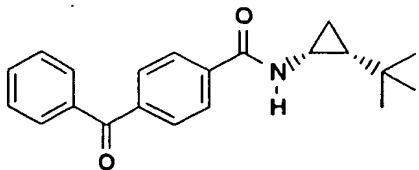
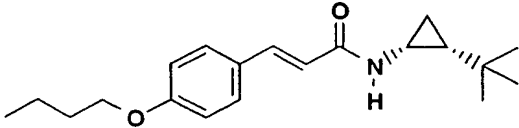
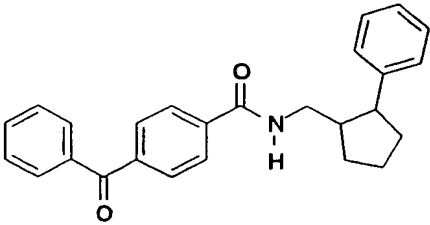
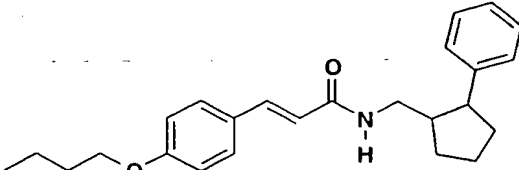
Example #	Structure	Characterization
8		$C_{19}H_{23}NO$: mp 126-127°C. Analysis calculated: C, 81.00; H, 8.24; N, 4.98. Found: C, 81.17; H, 8.19; N, 4.93.
9		$C_{17}H_{24}N_2O$: m/e = 272.
10		$C_{20}H_{23}NO_2$: m/e = 309.
10a	 ca. 2:1 cis:trans mixture	ESI (m/z) 398 Rf (silica, 25% EtOAc/hex) 0.21.
10b		ESI (m/z) 338 Rf (silica, 25% EtOAc/hex) 0.24.
10c		mp 56.0-57.0°C ESI (m/z) 358

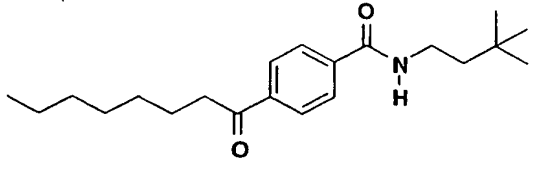
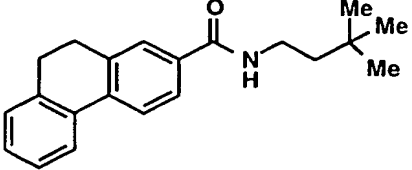
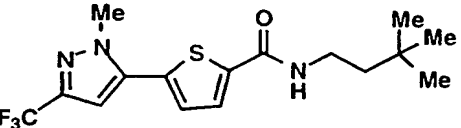
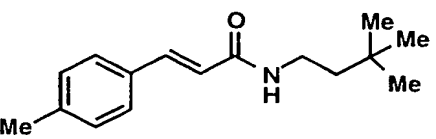
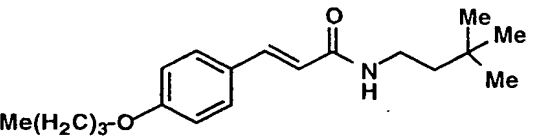
Example #	Structure	Characterization
10d		mp 62.5-64.0°C ESI (m/z) 332
10e		mp 50.5-51.5°C ESI (m/z) 332
10f	 ca. 2:1 mixture of isomers	mp 109.5-113.0°C ESI (m/z) 336
10g	 ca. 1.4:1 mixture of isomers	mp 133.0-133.5°C ESI (m/z) 350
10h		C ₂₂ H ₂₉ NO ₂ : mp 161.0-163.0°C. Analysis calculated: C, 81.69; H, 9.04; N, 4.33. Found: C, 81.65; H, 8.91; N, 4.13.

Example #	Structure	Characterization
10i		$C_{20}H_{21}NO_2 \cdot 0.27$ H_2O : mp 96.9-97.0°C. Analysis calculated: C, 76.94; H, 6.95; N, 4.49. Found: C, 76.94; H, 6.90; N, 4.59.
10j		ESI (m/z) 362 R_f (silica, 25% EtOAc/hex) 0.20.
10k		$C_{20}H_{21}NO_2 \cdot 0.16$ H_2O : mp 96.0-97.0°C. Analysis calculated: C, 77.40; H, 6.93; N, 4.51. Found: C, 77.40; H, 6.66; N, 4.47.
10l	 60:40 isomer mixture	$C_{22}H_{25}NO_2$: mp 106.0-107.5°C. Analysis calculated: C, 78.77; H, 7.51; N, 4.18. Found: C, 78.53; H, 7.51; N, 3.89.

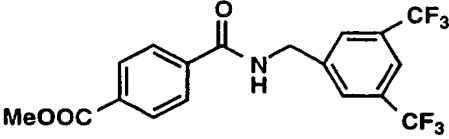
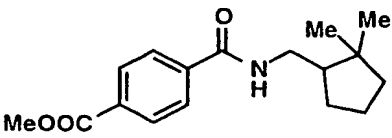
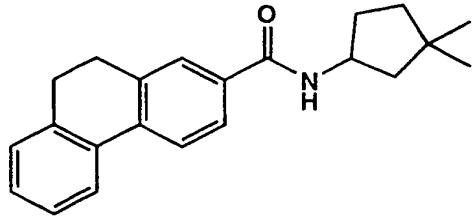
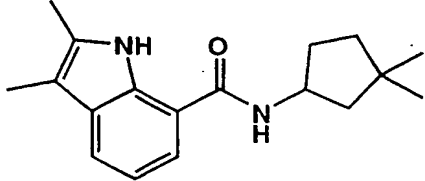
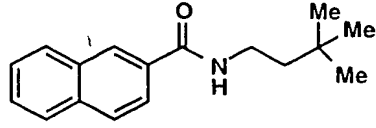
Example #	Structure	Characterization
10l'	 <p>60:40 isomer mixture</p>	<p>$C_{21}H_{23}NO_2$: mp 103.5-106.0°C.</p> <p>Analysis calculated: C, 78.47; H, 7.21; N, 4.36.</p> <p>Found: C, 78.32; H, 7.20; N, 4.29.</p>
10m		<p>mp 58.0-59.0°C</p> <p>ESI (m/z) 356.</p>
10m'		<p>$C_{19}H_{27}NO_2 \cdot 0.42 H_2O$: mp 104.0-107.5°C.</p> <p>Analysis calculated: C, 73.86; H, 9.08; N, 4.53.</p> <p>Found: C, 73.86; H, 9.26; N, 4.36.</p>
10n	 <p>60:40 mixture of isomers</p>	<p>$C_{20}H_{29}NO_2 \cdot 0.13 H_2O$: mp 107.0-110.0°C.</p> <p>Analysis calculated: C, 75.59; H, 9.28; N, 4.41.</p> <p>Found: C, 75.59; H, 9.51; N, 4.79.</p>

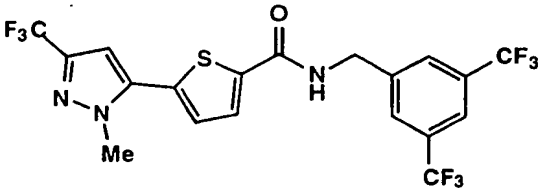
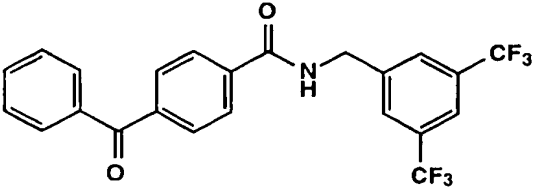
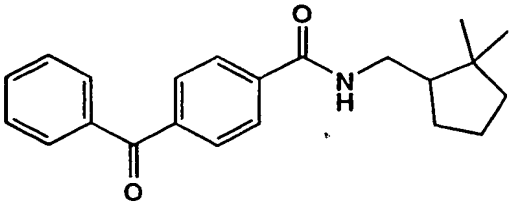
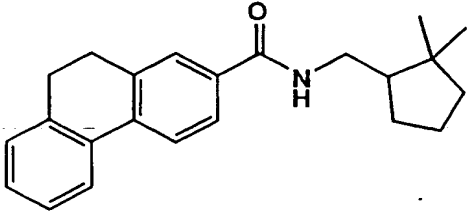
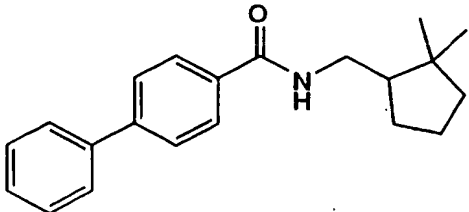
Example #	Structure	Characterization
10o	 <p>50:50 mixture of isomers</p>	$C_{21}H_{31}NO_2$: mp 90.0-92.5°C. Analysis calculated: C, 76.55; H, 9.48; N, 4.25. Found: C, 76.80; H, 9.73; N, 4.04.
10p		$C_{20}H_{29}NO_2 \cdot 0.18 H_2O$: mp 115.5-118.5°C. Analysis calculated: C, 75.38; H, 9.29; N, 4.40. Found: C, 75.38; H, 9.40; N, 4.65.
10q		$C_{19}H_{28}NO_2$: mp 88.5-90.0°C. Analysis calculated: C, 75.46; H, 9.33; N, 4.63. Found: C, 75.72; H, 9.19; N, 4.44.
10r		mp 85.5-87.0°C ESI (m/z) 322

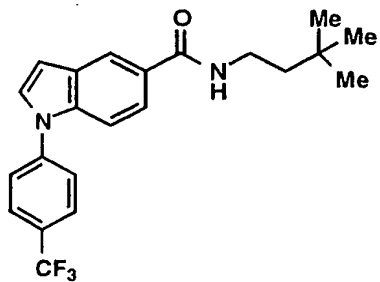
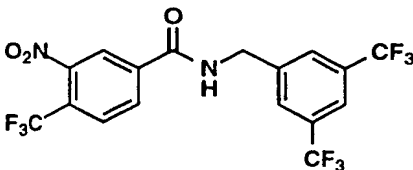
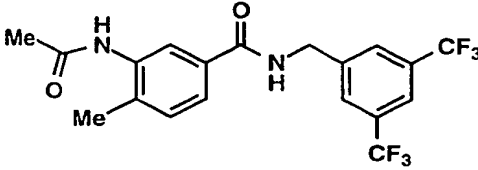
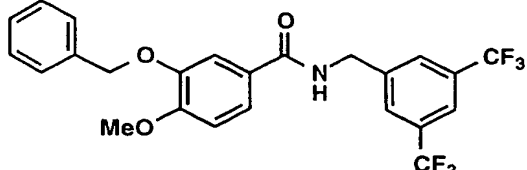
Example #	Structure	Characterization
10s		mp 122.0-125.0°C ESI (m/z) 322
10t		mp 151.0-153.0°C ESI (m/z) 316
10u	 <p>2:1 cis:trans mixture of isomers</p>	$C_{26}H_{25}NO_2 \cdot 0.15$ H ₂ O: ESI (m/z) 384. Analysis calculated: C, 80.85; H, 6.60; N, 3.63. Found: C, 80.85; H, 6.33; N, 3.52.
10v	 <p>2:1 cis:trans mixture of isomers</p>	$C_{25}H_{31}NO_2 \cdot 0.19$ H ₂ O: ESI (m/z) 378.. Analysis calculated: C, 78.83; H, 8.30; N, 3.68. Found: C, 78.83; H, 8.50; N, 3.57.

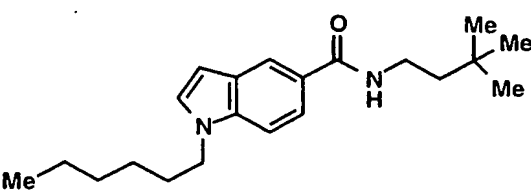
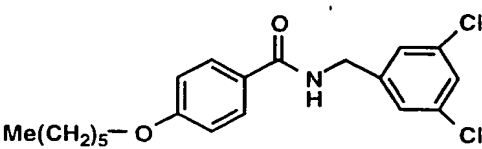
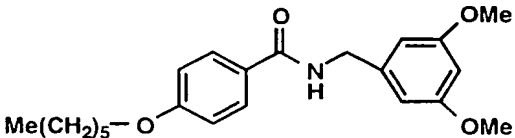
Example #	Structure	Characterization
10w		$C_{21}H_{33}NO_2$: mp 97.0-99.0°C. Analysis calculated: C, 76.09; H, 10.03; N, 4.23. Found: C, 75.81; H, 10.23; N, 4.13.
11		$C_{21}H_{25}NO$: m/e = 307.
12		$C_{16}H_{20}F_3N_3OS$: m/e = 359.
13		$C_{16}H_{23}NO$: m/e = 245.
14		$C_{19}H_{29}NO_2$: mp 82-84°C. Analysis calculated: C, 75.21; H, 9.63; N, 4.62. Found: C, 75.34; H, 9.44; N, 4.43.

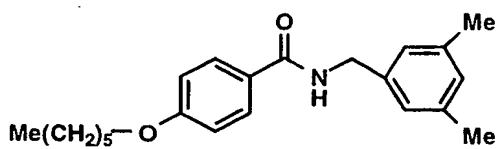
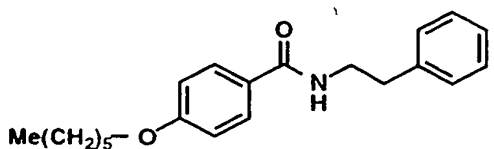
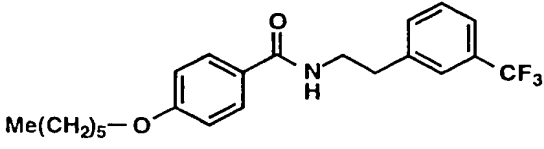
Example #	Structure	Characterization
14a		$C_{20}H_{29}NO_2$; $m/z = 316$ (M+H)
14b		$C_{22}H_{21}F_6NO_2$; $m/z = 446$ (M+H)
14c		MW $m/z = 304$ (M+H)
14d		$C_{21}H_{31}F_6NO_2$; $m/z = 330$ (M+H)
15		$C_{19}H_{30}N_2O_2 \cdot 0.25 H_2O$; mp $221^\circ C$. Analysis calculated: C, 70.67; H, 9.52; N, 8.67. Found: C, 70.67; H, 9.58; N, 8.57.

Example #	Structure	Characterization
16		$C_{18}H_{13}F_6NO_3$: mp 149-150°C. Analysis calculated: C, 54.05; H, 3.35; H, 3.50. Found: C, 53.60; H, 3.07; N, 3.48.
17		$C_{17}H_{23}NO_3$: mp 80-81°C. Analysis calculated: C, 70.56; H, 8.01; N, 4.84. Found: C, 70.27; H, 8.06; N, 4.74.
17a		$C_{22}H_{25}NO$: m/z = 320 (M+H)
17b		$C_{18}H_{24}N_2O$: m/z = 285 (M+H)
18		$C_{17}H_{21}NO$: m/e = 255.18

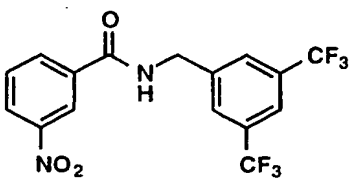
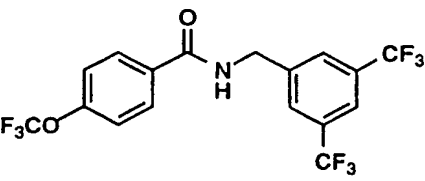
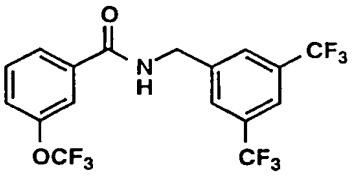
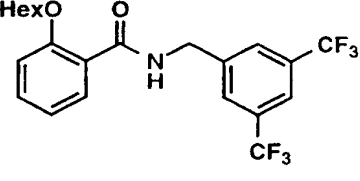
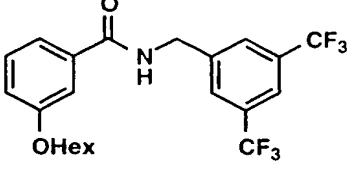
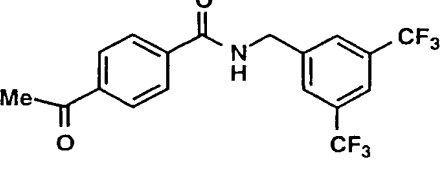
Example #	Structure	Characterization
18a		$C_{19}H_{12}F_9N_3OS$: $m/z = 501 (M+H)$
18b		$C_{23}H_{15}F_6NO_2$: $m/z = 452 (M+H)$
18c		$C_{22}H_{25}NO_2$: $m/z = 336 (M+H)$
18d		$C_{23}H_{27}NO$: $m/z = 334 (M+H)$
18e		$C_{21}H_{25}NO$: $m/z = 308 (M+H)$

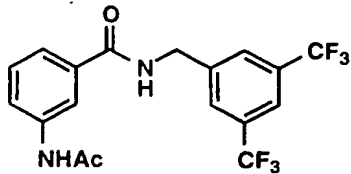
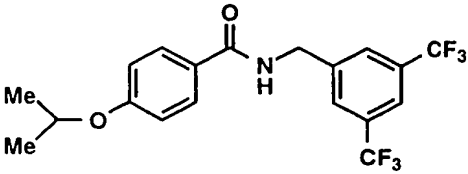
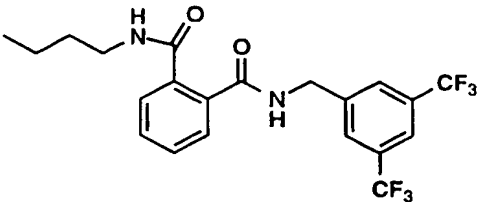
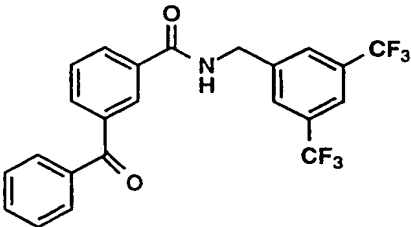
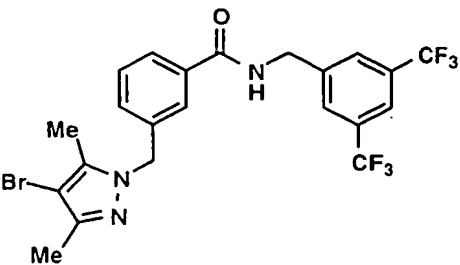
Example #	Structure	Characterization
19		$C_{22}H_{23}F_3N_2O$: mp 135-137°C. Analysis calculated: C, 68.02; H, 5.97; N, 7.21; F, 14.67. Found: C, 67.78; H, 5.71; N, 7.02; F, 14.67.
19a		$C_{17}H_9F_9N_2O_3$: m/z 460
19b		$C_{19}H_{16}F_6N_2O_2$: m/z 418
19c		$C_{24}H_{19}F_6NO_3$: m/z 483

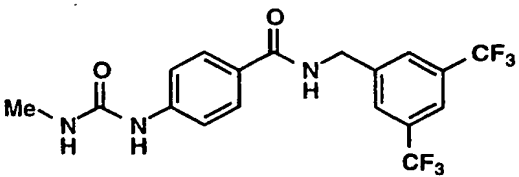
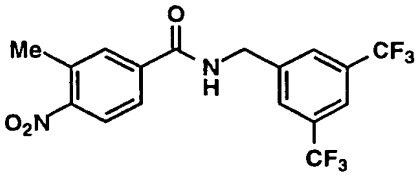
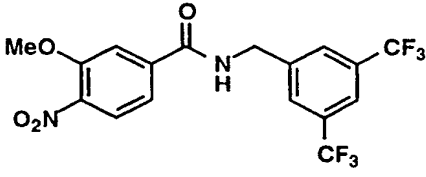
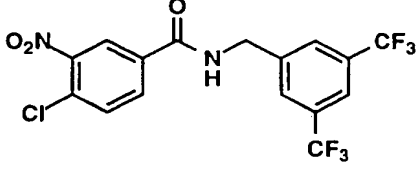
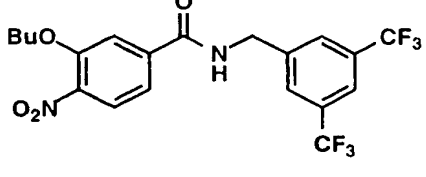
Example #	Structure	Characterization
20		$C_{21}H_{32}N_2O \cdot 0.32 H_2O$: mp 86-87°C. Analysis calculated: C, 75.46; H, 9.84; N, 8.38. Found: C, 75.44; H, 9.67; N, 8.27.
20a		$C_{20}H_{23}Cl_2NO_2$: mp 84-94°C. Analysis calculated: C, 63.16; H, 6.10; N, 3.68; Cl, 18.64. Found: C, 63.19; H, 5.99; N, 3.55; Cl, 18.58.
20b		$C_{22}H_{29}NO_4$: mp 80-83°C. Analysis calculated: C, 71.13; H, 7.87; N, 3.77. Found: C, 71.12; H, 7.86; N, 3.67.

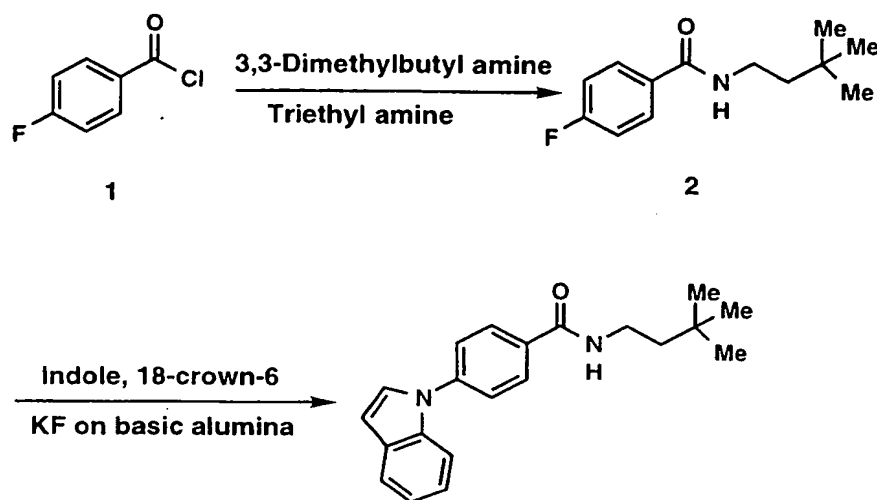
Example #	Structure	Characterization
20c	 <chem>CCCCCOc1ccc(cc1)C(=O)NCCc2cc(C)c(C)cc2</chem>	$C_{22}H_{29}NO_2$: mp 116-118°C. Analysis calculated: C, 77.84; H, 8.61; N, 4.13. Found: C, 78.05; H, 8.58; N, 4.10.
20d	 <chem>CCCCCOc1ccc(cc1)C(=O)NCCc2ccccc2</chem>	$C_{22}H_{29}NO_2$: mp 109-111°C. Analysis calculated: C, 77.50; H, 8.36; N, 4.30. Found: C, 77.69; H, 8.47; N, 4.31.
20e	 <chem>CCCCCOc1ccc(cc1)C(=O)NCCc2cc(C(F)(F)F)ccc2</chem>	$C_{22}H_{26}F_3NO_2$: mp 86-87°C. Analysis calculated: C, 67.16; H, 6.66; N, 3.56; F, 14.49. Found: C, 67.11; H, 6.62; N, 3.54; F, 14.71.

Example #	Structure	Characterization
20f		$C_{23}H_{25}F_6NO_2$: mp 86-87°C. Analysis calculated for 0.1 mole hexane: C, 60.30; H, 5.66; N, 2.98; F, 24.25. Found: C, 59.91; H, 5.33; N, 3.00; F, 23.88.
20g		$C_{17}H_{10}F_6N_2O$: m/z 372
20h		$C_{16}H_{10}ClF_6NO$: m/z 381
20i		$C_{17}H_{10}F_9NO$: m/z 415
20j		$C_{18}H_{16}F_6N_2O$: m/z 390

Example #	Structure	Characterization
20k		$C_{16}H_{10}F_6N_2O_3$ m/z 392
20l		$C_{17}H_{10}F_9NO_2$ m/z 431
20m		$C_{17}H_{10}F_9NO_2$ m/z 431
20n		$C_{22}H_{23}F_6NO_2$ m/z 447
20o		$C_{22}H_{23}F_6NO_2$ m/z 447
20p		$C_{18}H_{13}F_6NO_2$ m/z 389

Example #	Structure	Characterization
20q		$C_{18}H_{14}F_6N_2O_2$: m/z 404
20r		$C_{19}H_{17}F_6NO_2$: m/z 405
20s		$C_{21}H_{20}F_6N_2O_2$: m/z 446
20t		$C_{23}H_{15}F_6NO_2$: m/z 451
20u		$C_{22}H_{18}BrF_6N_3O$: m/z 534

Example #	Structure	Characterization
20v		$C_{18}H_{15}F_6N_3O_2$: m/z 419
20w		$C_{17}H_{12}F_6N_2O_3$: m/z 406
20x		$C_{17}H_{12}F_6N_2O_4$: m/z 422
20y		$C_{16}H_9ClF_6N_2O_3$: m/z 426
20z		$C_{20}H_{18}F_6N_2O_4$: m/z 464

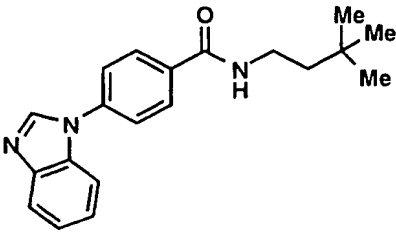
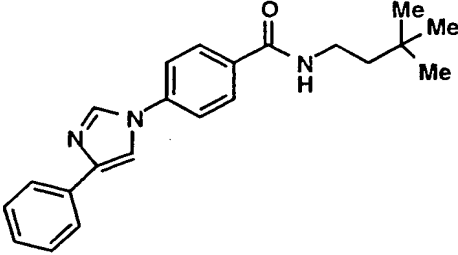
Example 21**N-(3,3-Dimethylbutyl)-4-(1H-indol-1-yl)benzamide****A. N-(3,3-Dimethylbutyl)-4-fluorobenzamide**

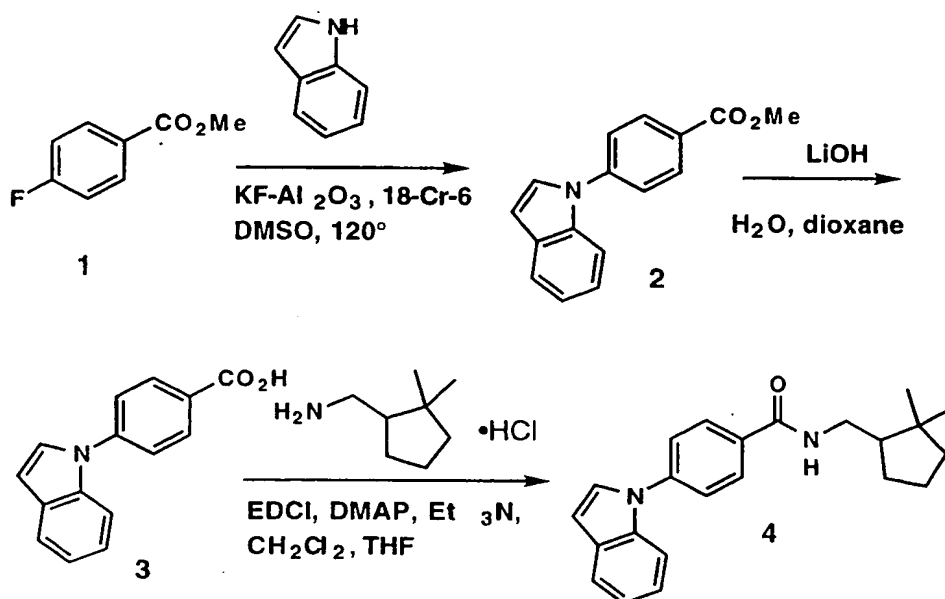
This compound was prepared from 4-fluorobenzoyl chloride (1) and 3,3-dimethylbutyl amine by the same procedure as described for the title compound of Example 1, part C.

B. N-(3,3-Dimethylbutyl)-4-(1H-indol-1-yl)benzamide

Compound 2 (186 mg, 0.83 mmol) and indole (127 mg, 1.08 mmol) were dissolved in dimethylsulfoxide (8 mL). 18-Crown-6 (66 mg, 0.25 mmol) and 37% by weight KF on basic alumina were added and the reaction mixture was heated at 120°C for 4 days. The reaction was cooled to room temperature, diluted with ethyl acetate and washed with brine. The organic layer was dried over magnesium sulfate and the solvent was removed to yield an orange oil. Purification by flash chromatography on silica gel eluting with 10% ethyl acetate in hexane provided a tan solid (180 mg, 67%): mp 51-54°C. Analysis calculated for C₂₁H₂₄N₂O: C, 78.72; H, 7.55; N, 8.74. Found: C, 78.59; H, 7.41; N, 8.69.

Using methodology analogous to that described for the title compound of Example 21, the compounds of Examples 22 and 23 were prepared.

Example #	Structure	Characterization
22		$C_{20}H_{23}N_3O \cdot 0.8H_2O$: mp 134-135°C. Analysis calculated: C, 71.53; H, 7.38; N, 12.51. Found: C71.42; H, 6.98; N, 12.43.
23		$C_{22}H_{25}N_3O \cdot 0.55H_2O$: mp 170-171°C. Analysis calculated: C, 73.94; H, 7.36; N, 11.76. Found: C, 73.94; H, 6.97; N, 11.74.

Example 23a**N-(3,3-Dimethylcyclopentyl)-4-(1H-indol-1-yl)benzamide****A. 4-(1H-Indol-1-yl)benzoic acid methyl ester**

Ethyl-4-fluorobenzoate (4 mL, 27.3 mmol) and indole (3.51 g, 30 mmol) were dissolved in 50 mL of dimethylsulfoxide (DMSO). Potassium fluoride (37% on basic alumina, 12.8 g) and 18-crown-6 (720 mg, 2.73 mmol) were added and the mixture was heated to 120°C for 24 hours. The reaction was diluted with ethyl acetate and washed with

hydrochloric acid (1.0 M, aq.), sodium bicarbonate (sat'd., aq.) and sodium chloride (sat'd., aq.). The organic layer was dried over magnesium sulfate, filtered and the solvent removed. Purification by flash chromatography on silica gel eluted with 5% ethyl acetate, hexane provided 1.45 g (20%) of a white solid

B. 4-(1H-Indol-1-yl)benzoic acid

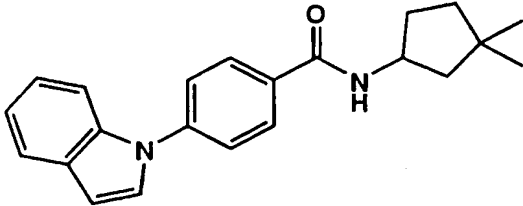
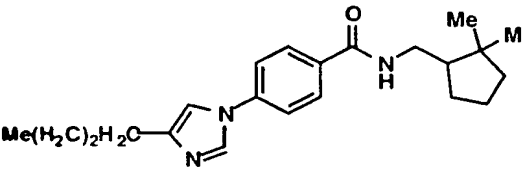
The ester (2, 1.4 g, 5.23 mmol) was dissolved in 45 mL of dioxane. Lithium hydroxide (15.8 mL, 15.8 mmol) was added and the mixture was stirred for 4 hours. The reaction was diluted with water and washed
5 with ethyl acetate. The aqueous phase was acidified with HCl and extracted with ethyl acetate. The organic layer was dried over magnesium sulfate, filtered and the solvent removed to provide 1.04 g (83%) of a pale yellow solid. mp 222-224°C.

10 C. N-(3,3-Dimethylcyclopentyl)-4-(1H-indol-1-yl)benzamide

The acid (3, 0.20 mmol) and the amine compound 3 from Example 1 (0.22 mmol) were dissolved in 1 mL of THF and triethylamine (0.22 mmol) and 4-dimethylaminopyridine (0.02 mmol) were added. 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (1.6 mL, 0.15
15 M in dichloromethane, 0.24 mmol) was added and the mixture was allowed to stand for 48 hours. Purification by preparative HPLC (YMC ODS-A 5S 30 x 250 mm column, 50-90% methanol water with 0.10% TFA gradient over 20 minutes, 25 mL/minute flow) provided 50 mg (72%) of a white solid. $C_{23}H_{26}N_2O$: $m/z = 347$ (M+H).

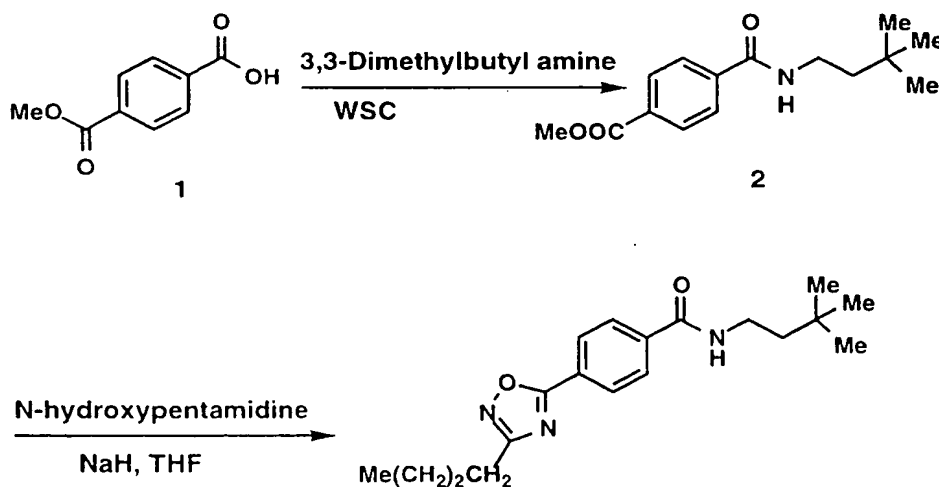
20

The following compounds can be prepared using methodology analogous to that described for the title compound of Example 23a above.

Example #	Structure	Characterization
23b		$C_{22}H_{24}N_2O$: $m/z = 333$ (M+H)
23c		$C_{22}H_{31}N_3O \cdot 0.67 H_2O$: Analysis calculated: C, 72.29; H, 8.92; N, 11.5. Found: C, 72.29; H, 9.11; N, 11.34.

Example 24

5 4-(3-Butyl-1,2,4-oxadiazol-5-yl)-N-(3,3-dimethylbutyl)benzamide



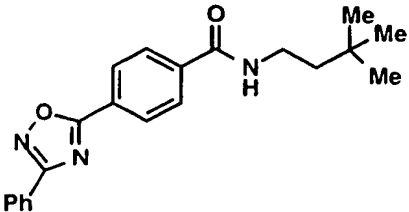
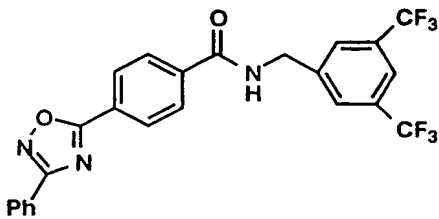
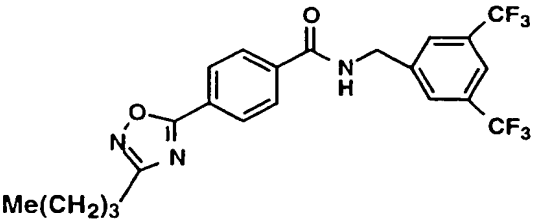
A. N-(3,3-Dimethylbutyl)-4-fluorobenzamide

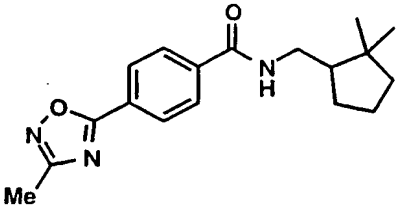
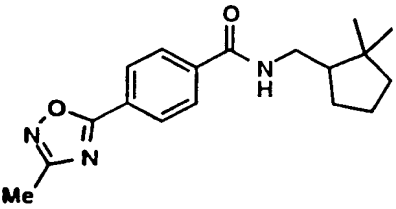
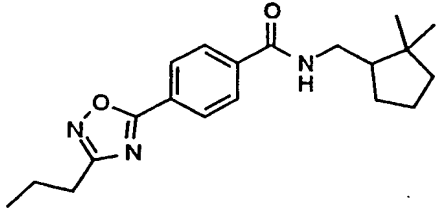
A solution of *mono*-methylterephthalate (1) (260 mg, 1.44 mmol) in DMF (3 mL) under argon at room temperature was treated with 3,3-dimethylbutylamine (145 mg, 1.43 mmol), ethyl-3-(3-dimethylamino)propyl carbodiimide hydrochloride (420 mg, 1.41 mmol) and hydroxybenzotriazole monohydrate (192 mg, 1.42 mmol) and stirred overnight. The reaction mixture was diluted with ethyl acetate and washed with 10% citric acid, water, sodium bicarbonate, water and brine. The dried (anhydrous magnesium sulfate) organic fraction was concentrated and the residue was purified by flash chromatography on silica gel, eluting with ethyl acetate/hexanes (1:4) to give compound 2 (327 mg, mp 88-90°C).

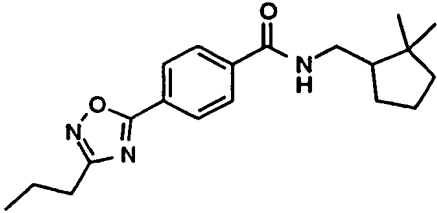
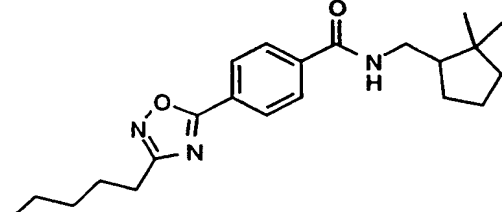
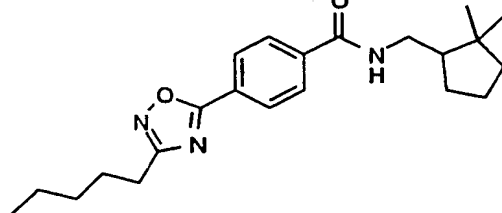
B. 4-(3-Butyl-1,2,4-oxadiazol-5-yl)-N-(3,3-dimethylbutyl)-benzamide

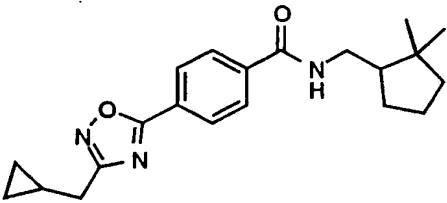
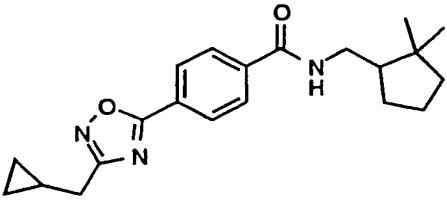
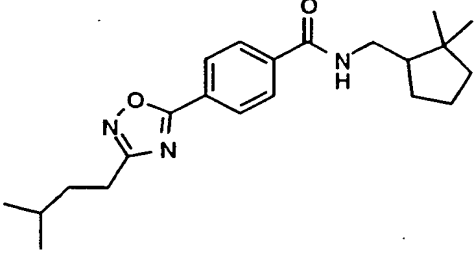
A solution of compound 2 (137 mg, 0.52 mmol) and N-hydroxypentamidine (75 mg, 0.65 mmol) in dimethylformamide (1.5 mL) under argon at 0-5°C was treated with sodium hydride (45 mg, 60%/m.o., 1.14 mmol) and allowed to stir at room temperature for 4 hours. The mixture, diluted with ethyl acetate, was washed with water and brine. The organic fraction was dried (anhydrous magnesium sulfate) and concentrated *in vacuo* to give a solid. Flash chromatography on silica gel, eluting with ethyl acetate/hexanes (1:4) gave the title compound (140 mg, 81%): mp 95-97°C. Analysis calculated for C₁₉H₂₇N₃O₂: C, 69.27; H, 8.26; N, 12.75. Found: C, 69.01; H, 8.29; N, 12.50.

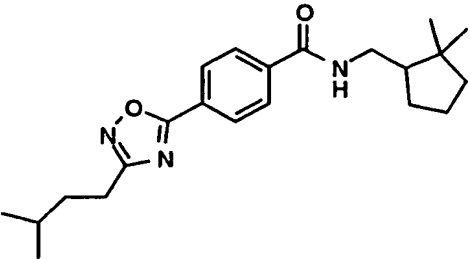
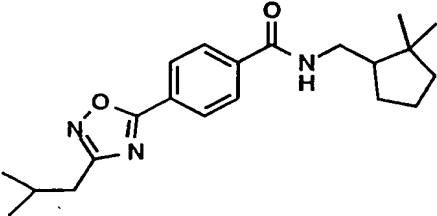
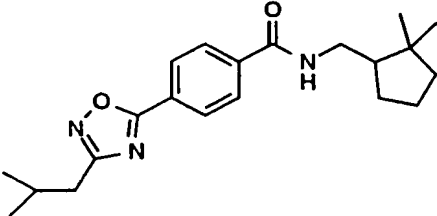
Using methodology analogous to that described for the title compound of Example 24, the compounds of Examples 25 to 29 were prepared.

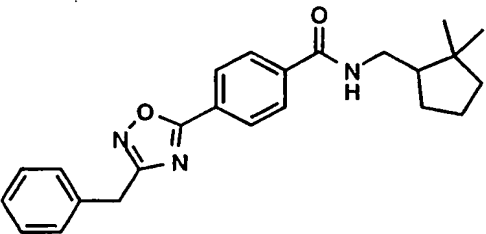
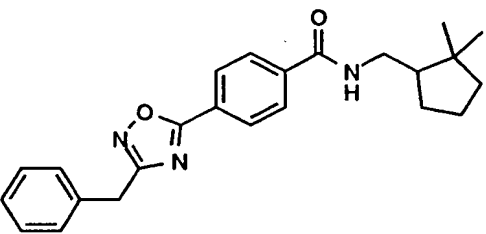
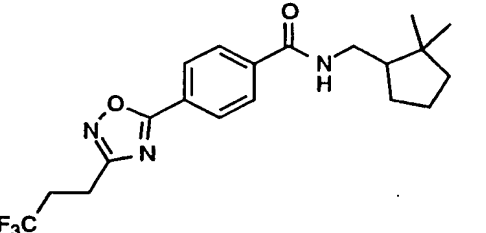
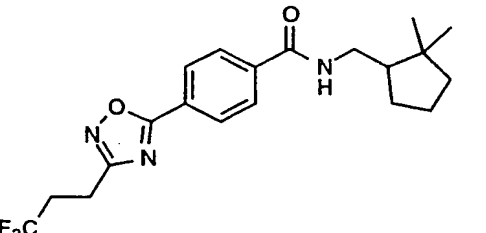
Example #	Structure	Characterization
25		$C_{21}H_{23}N_3O_2$: mp 169-170°C. Analysis calculated: C, 72.18; H, 6.63; N, 12.03. Found: C, 71.92; H, 6.63; N, 11.63.
26		$C_{24}H_{15}F_6N_3O_2$: mp 209-210°C. Analysis calculated: C, 58.66; H, 3.08; N, 8.55; F, 23.20. Found: C, 58.52; H, 3.08; N, 8.51; F, 22.85.
27		$C_{22}H_{19}F_6N_3O_2$: mp, 156-157°C. Analysis calculated: C, 56.56; H, 4.15; N, 8.99; F, 23.45. Found: C, 56.23; H, 3.88; N, 8.89; F, 23.28.

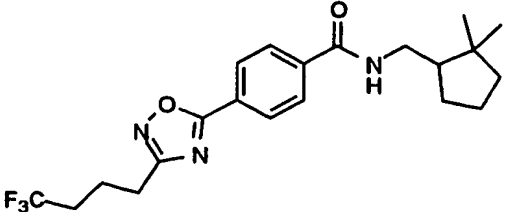
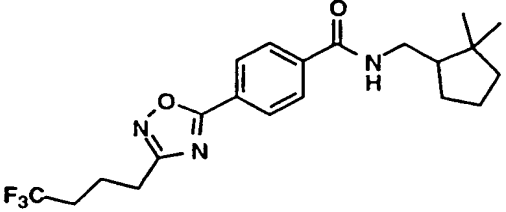
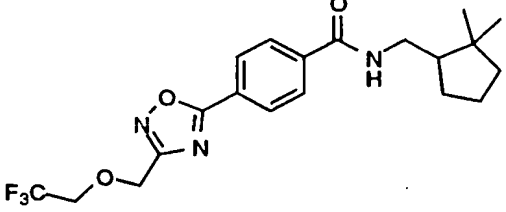
Example #	Structure	Characterization
27a		$C_{18}H_{23}N_3O_2$: mp 155-156°C. $[\alpha]_D = +22.8^\circ$ (c= 1.02, $CHCl_3$). Analysis calculated for 0.56 mole water: C, 66.83; H, 7.52; N, 12.99. Found: C, 66.82; H, 7.01; N, 12.57.
27b		$C_{18}H_{23}N_3O_2$: mp 155-156°C. $[\alpha]_D = -22.8^\circ$ (c= 1.00, $CHCl_3$). Analysis calculated: C, 68.98; H, 7.40; N, 13.41. Found: C, 68.78; H, 6.85; N, 13.10.
27c		$C_{20}H_{27}N_3O_2$: mp 88-90°C. $[\alpha]_D = +21.4^\circ$ (c= 0.49, $CHCl_3$). Analysis calculated: C, 70.35; H, 7.97; N, 12.31. Found: C, 70.09; H, 7.91; N, 12.17.

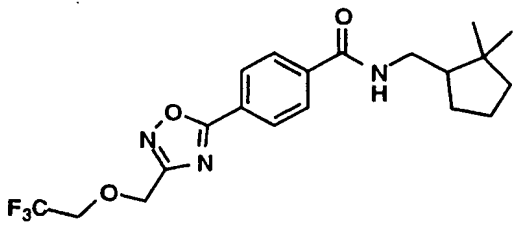
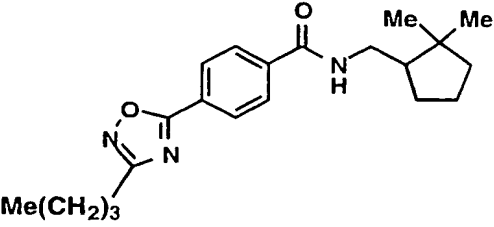
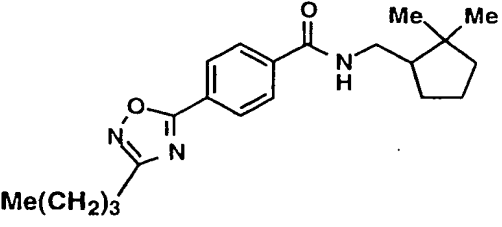
Example #	Structure	Characterization
27d		$C_{20}H_{27}N_3O_2$: mp 88-90°C. $[\alpha]_D = -21.6^\circ$ (c= 0.50, $CHCl_3$). Analysis calculated: C, 70.35; H, 7.97; N, 12.31. Found: C, 70.14; H, 7.84; N, 12.09.
27e		$C_{20}H_{27}N_3O_2$: mp 94-95°C. $[\alpha]_D = +18.8^\circ$ (c= 0.32, $CHCl_3$). Analysis calculated: C, 71.51; H, 8.46; N, 11.37. Found: C, 71.33; H, 8.49; N, 11.29.
27f		$C_{20}H_{27}N_3O_2$: mp 94-95°C. $[\alpha]_D = +19.5^\circ$ (c= 0.41, $CHCl_3$). Analysis calculated: C, 71.51; H, 8.46; N, 11.37. Found: C, 71.60; H, 8.47; N, 11.20.

Example #	Structure	Characterization
27g		$C_{21}H_{27}N_3O_2$: mp 116-117°C. $[\alpha]_D = +16.6^\circ$ (c= 0.35, $CHCl_3$). Analysis calculated: C, 71.36; H, 7.70; N, 11.89. Found: C, 71.57; H, 7.81; N, 11.76.
27h		$C_{20}H_{27}N_3O_2$: mp 94-95°C. $[\alpha]_D = -17.6^\circ$ (c= 0.33, $CHCl_3$). Analysis calculated: C, 71.51; H, 8.46; N, 11.37. Found: C, 71.29; H, 7.68; N, 11.78.
27i		$C_{22}H_{31}N_3O_2$: mp 92-93°C. $[\alpha]_D = +18.4^\circ$ (c= 0.25, $CHCl_3$). Analysis calculated: C, 71.51; H, 8.46; N, 11.37. Found: C, 71.37; H, 8.33; N, 11.14.

Example #	Structure	Characterization
27j		$C_{22}H_{31}N_3O_2$: mp 94-95°C. $[\alpha]_D = -17.0^\circ$ ($c = 0.20$, $CHCl_3$). Analysis calculated: C, 71.51; H, 8.46; N, 11.37. Found: C, 71.51; H, 8.58; N, 11.30.
27k		$C_{21}H_{29}N_3O_2$: mp 114-115°C. $[\alpha]_D = +18.1^\circ$ ($c = 0.43$, $CHCl_3$). Analysis calculated: C, 70.96; H, 8.22; N, 11.82. Found: C, 70.83; H, 8.30; N, 11.63.
27l		$C_{21}H_{29}N_3O_2$: mp 116-117°C. $[\alpha]_D = -18.0^\circ$ ($c = 0.20$, $CHCl_3$). Analysis calculated: C, 70.96; H, 8.22; N, 11.82. Found: C, 70.80; H, 8.22; N, 11.70.

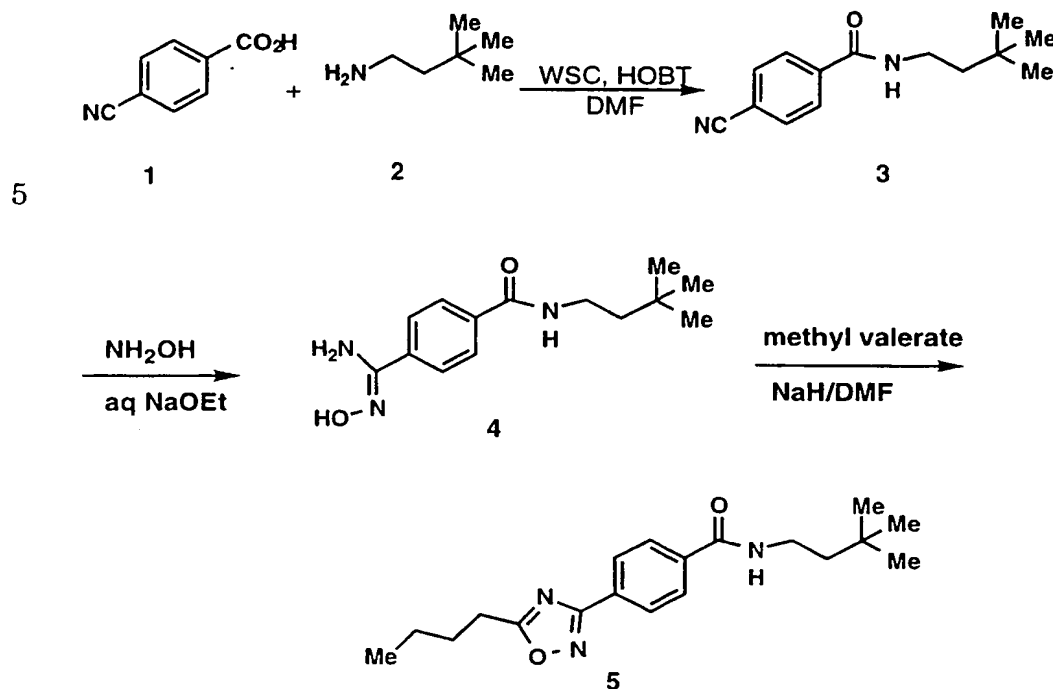
Example #	Structure	Characterization
27m		$C_{24}H_{27}N_3O_2 \cdot 0.16H_2O \cdot 0.03EtOAc$: mp 138-139°C. $[\alpha]_D = +15.1^\circ$ (c= 0.37, $CHCl_3$). Analysis calculated: C, 73.25; H, 7.02; N, 10.62. Found: C, 73.04; H, 6.96; N, 10.45.
27n		$C_{24}H_{27}N_3O_2$: mp 139-140°C. $[\alpha]_D = -15.6^\circ$ (c= 0.23, $CHCl_3$). Analysis calculated: C, 74.01; H, 6.99; N, 10.79. Found: C, 73.73; H, 6.77; N, 10.54.
27o		$C_{20}H_{24}F_3N_3O_2$ $[\alpha]_D = +11.8^\circ$ (c= 0.49, $CHCl_3$). m/s MW = 395.18
27p		$C_{20}H_{24}F_3N_3O_2$ $[\alpha]_D = -13.7^\circ$ (c= 0.90, $CHCl_3$). m/s MW = 395.18

Example #	Structure	Characterization
27q		$C_{21}H_{26}F_3N_3O_2$; mp 83-84°C. $[\alpha]_D = +15.6^\circ$ (c= 0.23 $CHCl_3$). Analysis calculated: C, 61.60; H, 6.40; N, 10.26; F, 13.92. Found: C, ; H, ; N, ; F, .
27r		$C_{21}H_{26}F_3N_3O_2$; mp 83-84°C. $[\alpha]_D = -15.0^\circ$ (c= 0.36, $CHCl_3$). Analysis calculated: C, 61.60; H, 6.40; N, 10.26; F, 13.92. Found: C, 61.58; H, 6.49; N, 10.19; F, 13.66.
27s		$C_{20}H_{24}F_3N_3O_3$; mp °C. $[\alpha]_D = +13.8^\circ$ (c= 0.24, $CHCl_3$). Analysis calculated: C, 58.39; H, 5.88; N, 10.21; F, 13.85. Found: C, ; H, ; N, ; F, .

Example #	Structure	Characterization
27t		$C_{20}H_{24}F_3N_3O_3$; mp 83-84.5°C. $[\alpha]_D = -17.5^\circ$ (c = 0.24 $CHCl_3$). Analysis calculated: C, 58.39; H, 5.88; N, 10.21; F, 13.85. Found: C, 58.52; H, 5.95; N, 10.16; F, 13.34.
28		$C_{21}H_{29}N_3O_2 \cdot 0.3$ hexanes: mp 74-76°C, $[\alpha]_D = +17.3^\circ$ (c = 0.11, $CHCl_3$). Analysis calculated: C, 71.81; H, 8.78; N, 11.02. Found: C, 71.85; H, 8.69; N, 11.09.
29		$C_{21}H_{29}N_3O_2 \cdot 0.3$ hexanes: mp 74-76°C, $[\alpha]_D = -20^\circ$ (c = 0.12, $CHCl_3$). Analysis calculated: C, 71.81; H, 8.78; N, 11.02. Found: C, 71.85; H, 8.69; N, 11.09.

Example 29a

4-(5-Butyl-1,2,4-oxadiazol-3-yl)-N-(3,3-dimethylbutyl)benzamide



A. 4-Cyano-N-(3,3-dimethylbutyl)benzamide

A solution of methyl 4-cyanobenzoic acid (1, 735 mg, 5.0 mmol) in 12 mL of DMF under argon at room temperature was treated 3,3-dimethylbutylamine (2, 505 mg, 5.0 mmol), ethyl-3-(3-dimethylamino)-propyl carbodiimide hydrochloride (1.5 g, 5.0 mmol) and hydroxybenzotriazole monohydrate (685 mg, 5.0 mmol) and stirred overnight. The reaction mixture was diluted with ethyl acetate and washed with 10 percent citric acid, water, dilute sodium bicarbonate, water and brine. The dried (anhydrous magnesium sulfate) organic fraction was concentrated to give 1.5 g. Flash chromatography on 150 mL of EM-60 silica gel, eluting with ethyl acetate/hexanes (1:4) gave the title compound (1.06 g, 92%), mp 154-157°C. Analysis calculated for C₁₄H₁₈N₂O: C, 73.01; H, 7.88; N, 12.16. Found: C, 73.00; H, 7.92; N, 12.17.

B. 4-[Amino(hydroxyimino)methyl]-N-(3,3-dimethylbutyl)benzamide

A slurry of hydroxylamine hydrochloride (100 mg, 1.4 mmole) in water (0.1 mL) at room temperature was treated with sodium ethoxide in ethanol (prepared from 33 mg sodium in 2 mL ethanol) and the title A compound and stirred for 48 hours. The mixture was diluted with a small amount of ethanol and filtered to remove solids. The filtrate was concentrated, redissolved in ethanol, cooled in ice and filtered. The filtrate was concentrated and dried under high vacuum over P_2O_5 to give 348 mg hydroxyamidine product 4. m/z MH^+ @ 264, MW = 263.

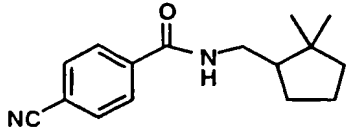
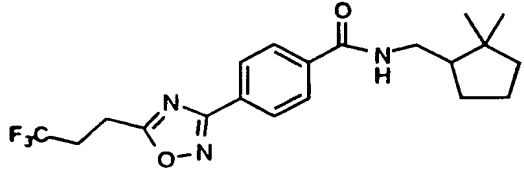
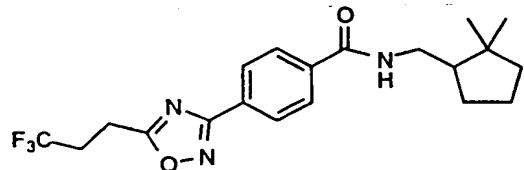
C. 4-(5-Butyl-1,2,4-oxadiazol-3-yl)-N-(3,3-dimethylbutyl)-benzamide

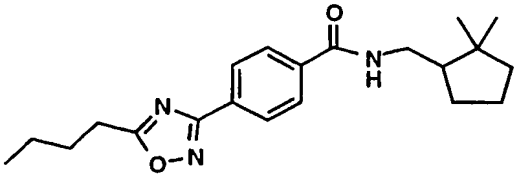
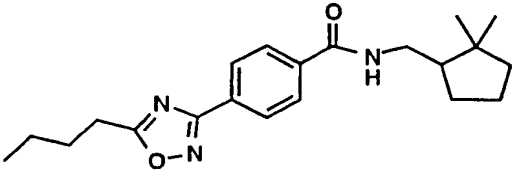
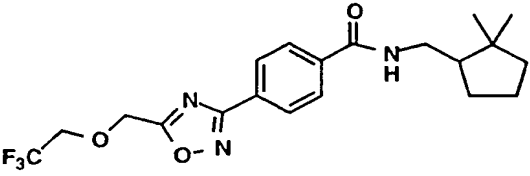
A solution of the title B compound (200 mg, 0.76 mmol) and methyl valerate (90 mg, 0.78 mmol) in 2 mL of dimethylformamide under argon at 0-5°C was treated with sodium hydride (33 mg, 60 %/m.o., 0.82 mmol) and allowed to stir at room temperature overnight. The mixture, diluted with ethyl acetate, was washed with water and brine. The organic fraction was dried (anhydrous magnesium sulfate) and concentrated. Flash chromatography on 150 mL of EM-60 silica gel, eluting with ethyl acetate/hexanes (1:4) gave the desired compound (145 mg, 58 %), mp 78-80°C. Analysis calculated for $C_{19}H_{27}N_3O_2$: C, 69.27; H, 8.26; N, 12.75.

Found:

C, 69.14; H, 8.37; N, 12.71.

Using the above methodology, the following compounds were prepared.

Example #	Structure	Characterization
29b		$C_{16}H_{20}N_2O$: mp 156-158.5°C. Analysis calculated: C, 74.97; H, 7.86; N, 10.93;. Found: C, 74.87; H, 7.99; N, 10.79.
29k		$C_{20}H_{24}F_3N_3O_2$: mp 116-118°C. $[\alpha]_D = +16.8^\circ$ (c= 0.88 $CHCl_3$). Analysis calculated: C, 60.75; H, 6.12; N, 10.63; F, 14.41. Found: C, 60.57; H, 6.22; N, 10.48; F, 14.02.
29l		$C_{20}H_{24}F_3N_3O_2$: mp 116-118°C. $[\alpha]_D = -16.9^\circ$ (c= 0.96 $CHCl_3$). Analysis calculated: C, 60.75; H, 6.12; N, 10.63; F, 14.41. Found: C, 60.87; H, 6.21; N, 10.45; F, 14.10.

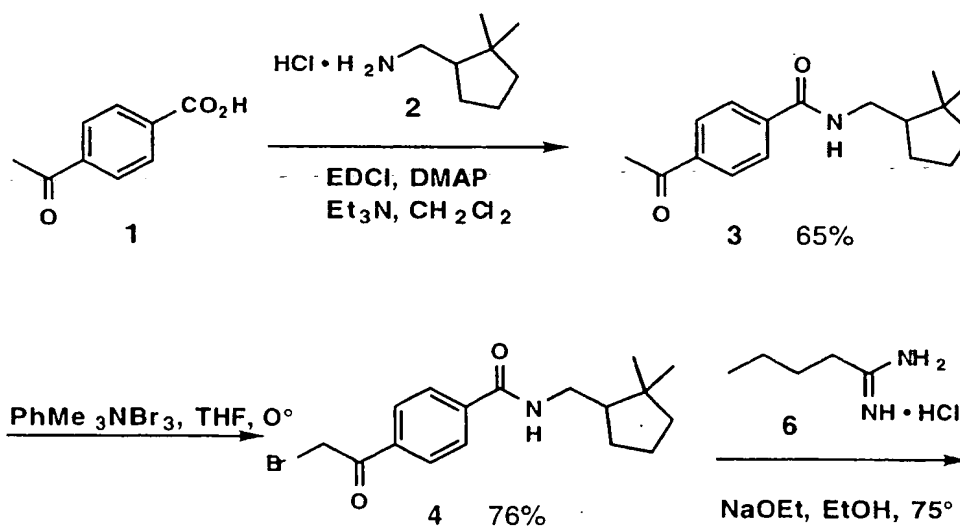
Example #	Structure	Characterization
29e		$C_{21}H_{29}N_3O_2 \cdot 0.1H_2O$ mp 86-88°C. $[\alpha]_D = +20.1^\circ$ (c = 0.63 $CHCl_3$). Analysis calculated: C, 70.60; H, 8.24; N, 11.76. Found: C, 70.57; H, 8.49; N, 11.26.
29f		$C_{21}H_{29}N_3O_2$: mp 86-88°C. $[\alpha]_D = +18.1^\circ$ (c = 0.63 $CHCl_3$). Analysis calculated: C, 70.96; H, 8.22; N, 11.82. Found: C, 70.76; H, 8.49; N, 11.57.
29g		$C_{20}H_{24}F_3N_3O_3$: mp 73-75°C. $[\alpha]_D = +17.6^\circ$ (c = 0.75 $CHCl_3$). Analysis calculated: C, 58.39; H, 5.88; N, 10.21; F, 13.85. Found: C, 58.51; H, 5.87; N, 10.02; F, 13.49.

Example #	Structure	Characterization
29h		$C_{20}H_{24}F_3N_3O_3$; mp 73-75° C. $[\alpha]_D = -17.2^\circ$ ($c = 0.89$ $CHCl_3$). Analysis calculated: C, 58.39; H, 5.88; N, 10.21; F, 13.85. Found: C, 57.98; H, 5.90; N, 9.87; F, 13.45.

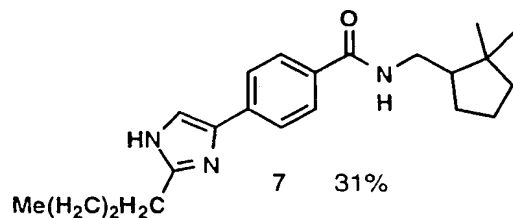
Example 29i

4-(2-Butyl-1H-imidazol-4-yl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide

5



10



A. 4-Acetyl-N-[(2,2-dimethylcyclopentyl)methyl]benzamide

4-Acetylbenzoic acid (200 mg, 1.20 mmol) and amine 2 (from Example 1, 199 mg, 1.22 mmol) were dissolved in 5 mL of dichloromethane and triethylamine (187 μ L, 1.34 mmol) and 4-dimethylaminopyridine (15 mg, 0.12 mmol) were added. 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (398 mg, 1.34 mmol) was added and the mixture was stirred for 18 hours. The reaction was diluted with ethyl acetate and washed with hydrochloric acid (1.0 M, aq.), sodium bicarbonate (sat'd., aq.) and sodium chloride (sat'd., aq.). The organic layer was dried over magnesium sulfate, filtered and the solvent removed to yield 216 mg (65%) of a white solid.

B. 4-(Bromoacetyl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide

The amide 3 (160 mg, 0.585 mmol) was dissolved in 3 mL of tetrahydrofuran (THF) and cooled to 0°C. Phenyltrimethyl-ammonium tribromide (220 mg, 0.585 mmol) was dissolved in 1 mL of THF and added dropwise. The reaction was allowed to warm to room temperature and stir for 22 hours. The mixture was quenched with water and extracted with ethyl acetate. The organic layer was dried over magnesium sulfate, filtered and the solvent removed to yield 230 mg of a tan solid. Purification by flash chromatography on silica gel eluted with 30% ethyl acetate, hexane provided 157 mg (76%) of a white solid.

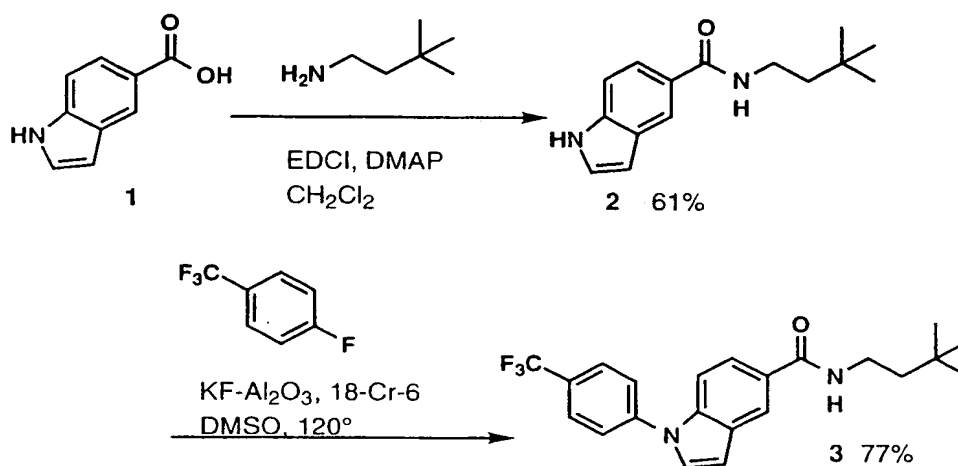
C. 4-(2-Butyl-1H-imidazol-4-yl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide

A solution of n-pentamidine hydrochloride (180 mg, 1.32 mmol) was dissolved in 1 mL of ethanol and sodium ethoxide (2 M in ethanol, 0.66 mL, 1.32 mmol) was added and the mixture was heated to 75°C. The title B compound (155 mg, 0.440 mmol) was dissolved in 1.5 mL of ethanol and added in small portions over 30 minutes. The reaction was stirred for 4 hours, quenched with sodium bicarbonate and the aqueous

layer was extracted with ethyl acetate. The organic extracts were dried over magnesium sulfate, filtered and the solvent removed to provide 173 mg of a brown solid. Purification by flash chromatography on silica gel eluted with 70% ethyl acetate, hexane provided 49 mg (31%) of a white solid, mp 98-105°C; Analysis calculated for $C_{22}H_{31}N_3O \cdot 0.59 H_2O$: C, 72.57; H, 8.91; N, 11.54. Found: C, 72.56; H, 9.00; N, 11.39.

Example 29j

N-(3,3-Dimethylbutyl)-1-[4-(trifluoromethyl)phenyl]-1H-indole-5-carboxamide



15 A. N-(3,3-Dimethylbutyl)-1H-indole-5-carboxamide

Indole-5-carboxylic acid (322 mg, 2.00 mmol), 3,3-dimethylbutylamine (296 μ L, 2.20 mmol) and 4-dimethyl-aminopyridine (24 mg, 0.20 mmol) were suspended in 10 mL of dichloromethane and 5 mL of N, N-dimethylformamide. 1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDCI, 653 mg, 2.2 mmol) was added and the mixture stirred for 16 hours. The reaction was diluted with ethyl acetate and washed with hydrochloric acid (1.0 M, aq.), sodium bicarbonate and brine. The organic layer was dried over magnesium sulfate, filtered and the solvent removed to yield a white solid (448 mg).

20 Recrystallization from hexane provided pale yellow needles which were dissolved in ether, filtered and the solvent removed to provide 300 mg

(61%) of an off white solid, mp 132-133°C; Analysis calculated for $C_{15}H_{20}N_2O \cdot 0.10 H_2O$: C, 73.20; H, 8.27; N, 11.38. Found: C, 73.17; H, 8.25; N, 11.20.

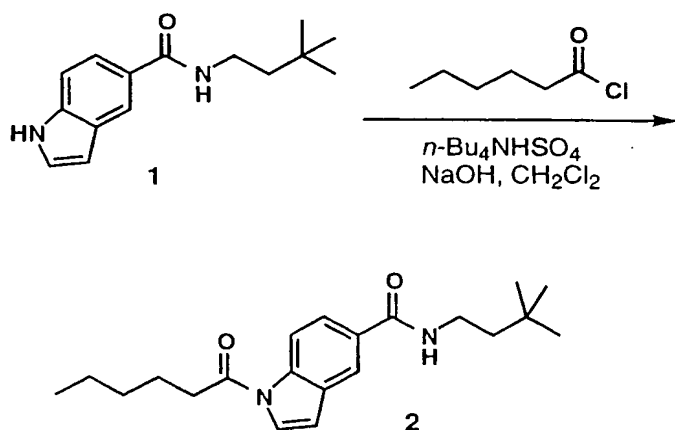
5 **B. N-(3,3-Dimethylbutyl)-1-[4-(trifluoromethyl)phenyl]-1H-indole-5-carboxamide**

The title A compound (40 mg, 0.16 mmol) and 4-fluorobenzotrifluoride (42 μ L, 0.33 mmol) were dissolved in 2 mL of dimethylsulfoxide. 18-Crown-6 (9 mg, 0.03 mmol) and 37% by weight KF
10 on basic alumina (51 mg, 0.33 mmol) were added and the mixture was heated to 120°C for 4 days. The reaction was cooled to room temperature, diluted with ethyl acetate and washed with sodium chloride (sat'd., aq.). The organic layer was dried over magnesium sulfate, filtered and the solvent removed to yield 57 mg of a yellow solid. Purification by flash
15 chromatography on silica gel eluted with 15% acetone, hexane provided 49 mg (77%) of a white solid. mp 135-137°C; Analysis calculated for $C_{22}H_{23}F_3N_2O$: C, 68.02; H, 5.97; N, 7.21; F, 14.67. Found: C, 67.78; H, 5.71; N, 7.02; F, 14.67.

20

Example 29k

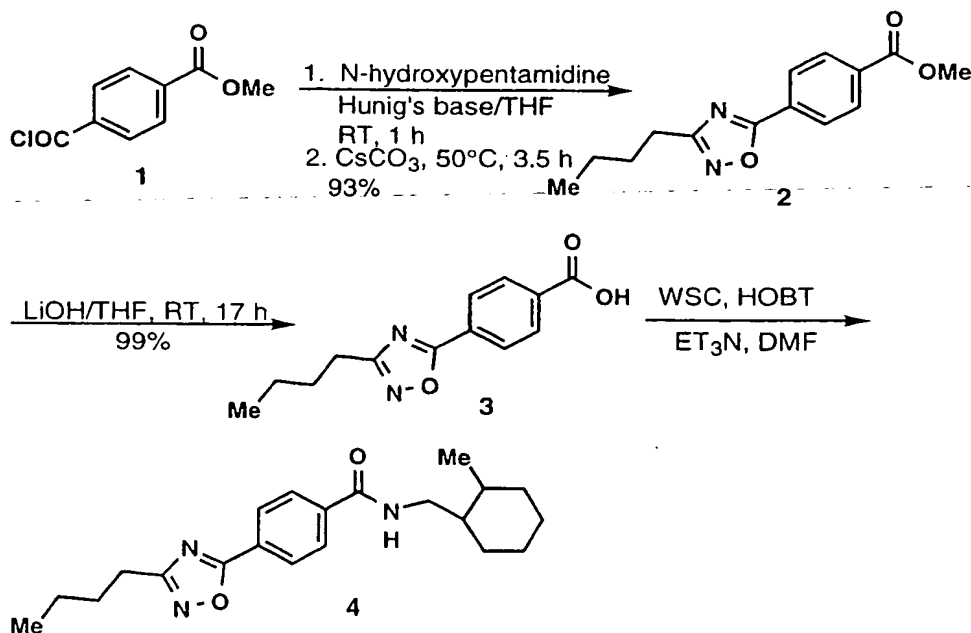
N-(3,3-Dimethylbutyl)-1-(1-oxohexyl)-1H-indole-5-carboxamide



The title A compound of example 29j (44 mg, 0.18 mmol) and tetrabutylammonium hydrogen sulfate (6 mg, 0.02 mmol) were dissolved in 1 mL of dichloromethane. Powdered sodium hydroxide (18 mg, 0.45 mmol) and hexanoyl chloride (38 μ L, 0.27 mmol) were added and the mixture was stirred for 2 hours. The reaction was diluted with ethyl acetate and washed with 1N hydrochloric acid, sodium bicarbonate and brine. The organic layer was dried over magnesium sulfate, filtered and the solvent removed to yield 58 mg of a white solid. Purification by flash chromatography on silica gel eluting with 20% acetone, hexane provided 52 mg (84%) of a white solid. mp 136-138°C; Analysis calculated for $C_{21}H_{30}N_2O_2$: C, 73.65; H, 8.83; N, 8.18. Found: C, 73.53; H, 8.94; N, 8.13.

Example 29k'

4-(3-Butyl-1,2,4-oxadiazol-5-yl)-N-[(2-methylcyclohexyl)methyl]-benzamide



A. 4-(3-Butyl-1,2,4-oxadiazol-5-yl)benzoic acid methyl ester

Diisopropylethylamine (1.1 ml, 6.0 mmol) was added to a solution of terephthalic acid, monomethyl ester chloride (1.0 g, 5.0 mmol) and N-hydroxypentamidine (0.70 g, 6.0 mmol) in THF (8 mL). After stirring at ambient temperature for 1 hour, cesium carbonate (3.6 g, 11 mmol) was added and the reaction was stirred at 50°C. After stirring at 50°C for 3.5 hours, the reaction was transferred to a separatory funnel with CH₂Cl₂/H₂O. The aqueous layer was acidified with 1N HCl to pH 1 and extracted with CH₂Cl₂ (3 x 40 mL). The combined organic layers were dried (MgSO₄) and concentrated *in vacuo*. Flash chromatography (silica, 50 mm dia, 10% EtOAc/hexane) afforded 1.2 g (93%) of methyl (3-butyl-1,2,4-oxadiazol-5-yl)benzoate.

B. 4-(3-Butyl-1,2,4-oxadiazol-5-yl)benzoic acid

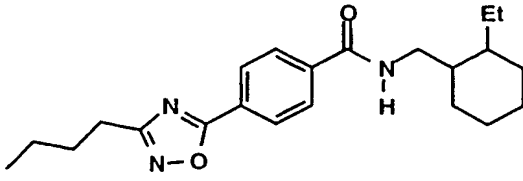
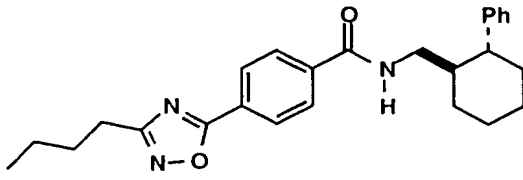
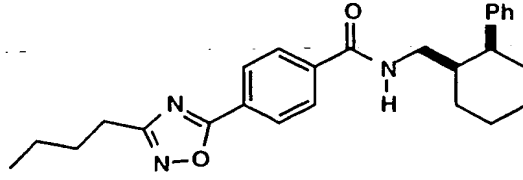
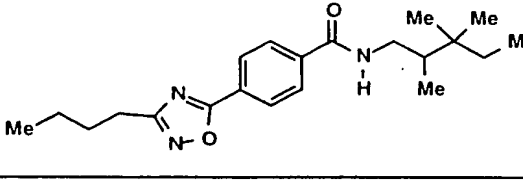
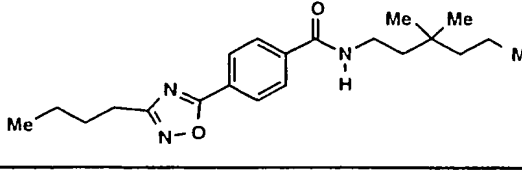
Lithium hydroxide (1.0 N, 9.8 ml, 9.8 mmol) was added to a stirring solution of the title A compound (1.2 g, 4.6 mmol) in THF (45 mL). After stirring at ambient temperature for 17 hours, the reaction was evaporated *in vacuo*. Water was added and the solution acidified with 1N HCl. The resulting solid was collected, washed with H₂O, and dried under high vacuum over P₂O₅ to afford 1.1 g (99%) of (3-butyl-1,2,4-oxadiazol-5-yl)benzoic acid.

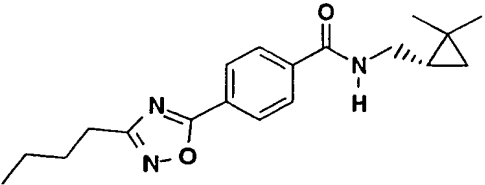
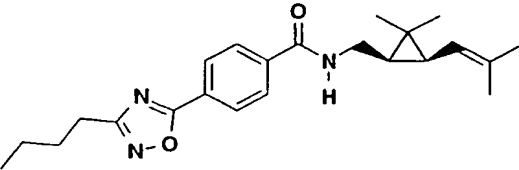
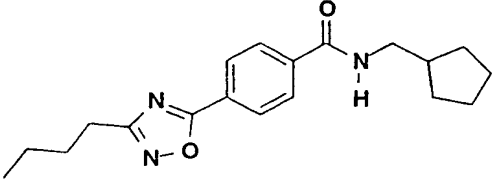
C. 4-(3-Butyl-1,2,4-oxadiazol-5-yl)-N-[(2-methylcyclohexyl)-methyl]benzamide

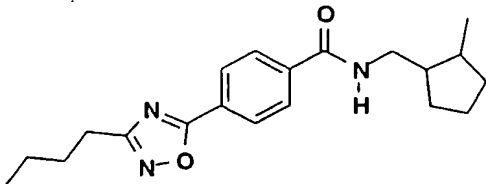
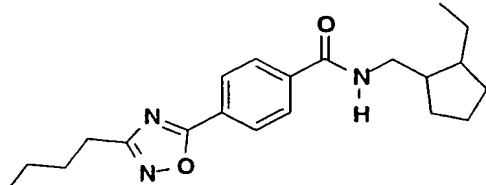
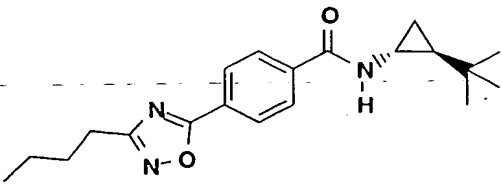
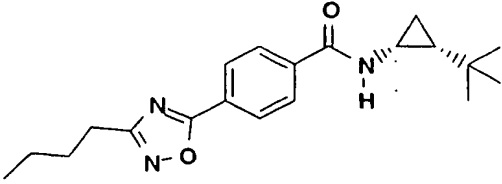
A solution of ((2-methylcyclohexyl)methyl)amine hydrochloride, prepared according to Example 1f, part B (104 mg, 0.633 mmol), title B compound (130 mg, 0.528 mmol), HOBT hydrate (86 mg), triethylamine (0.15 mL, 1.1 mmol), and 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (WSC, 121 mg, 0.633 mmol) in DMF (5 mL) was stirred for 17 hours. The reaction was diluted with ether (25 mL); washed with 0.3 N HCl (3 x 7 mL), and saturated NaHCO₃ (10 mL); dried (MgSO₄) and

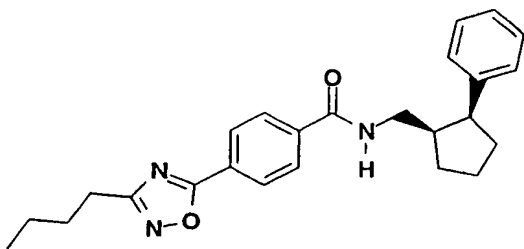
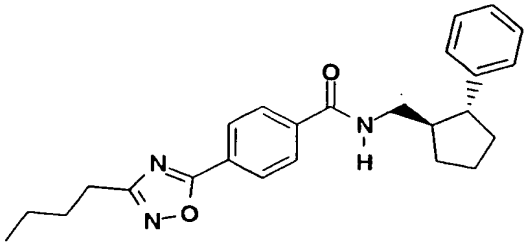
concentrated *in vacuo*. Purification by HPLC provided the title compound as a 1:1 mixture of isomers: MS: (ESI) m/z 356.

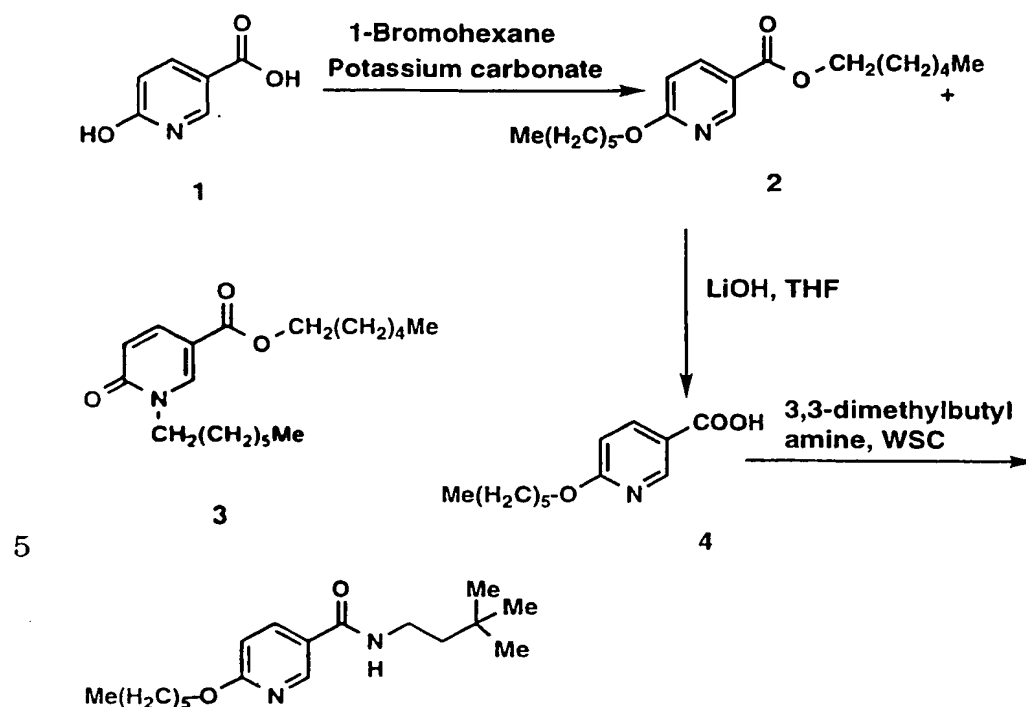
Using the procedure described for example 29k, the following
5 compounds were prepared.

Example #	Structure	Characterization
29l	 <p>ca 1:1 mixture of cis:trans</p>	ESI (m/z) 370 Rf (silica, 25% EtOAc/hex) 0.25.
29m	 <p>trans isomer</p>	mp 141.5-142.0°C ESI (m/z) 418
29n		mp 162.0-162.5°C ESI (m/z) 418
29o		mp 79.0-79.5°C ESI (m/z) 358
29p		mp 50.0-52.0°C ESI (m/z) 358

Example #	Structure	Characterization
29q		$C_{19}H_{25}N_3O_2$: mp 98.0-99.5°C. Analysis calculated: C, 69.70; H, 7.70; N, 12.83. Found: C, 69.70; H, 7.88; N, 12.68.
29r		$C_{23}H_{31}N_3O_2 \cdot 0.09 H_2O$: mp 59.0-60.0°C. Analysis calculated: C, 72.10; H, 8.20; N, 10.97. Found: C, 72.10; H, 8.49; N, 10.84.
29s		$C_{19}H_{25}N_3O_2$: mp 127.0-128.5°C. Analysis calculated: C, 69.70; H, 7.70; N, 12.83. Found: C, 69.53; H, 7.93; N, 12.64.

Example #	Structure	Characterization
29t	 <p>60:40 isomer mixture</p>	$C_{20}H_{27}N_3O_2$: mp 111.0-113.0°C. Analysis calculated: C, 70.35; H, 7.97; N, 12.31. Found: C, 70.24; H, 8.20; N, 12.27.
29u	 <p>50:50 isomer mixture</p>	$C_{21}H_{29}N_3O_2$: mp 121.0-124.5°C. Analysis calculated: C, 70.95; H, 8.22; N, 11.82. Found: C, 71.00; H, 8.40; N, 11.99.
29v		mp 105.0-107.0°C ESI (m/z) 342
29w		mp 132.0-133.5°C ESI (m/z) 342

Example #	Structure	Characterization
29x		$C_{25}H_{29}N_3O_2$: mp 112.0-115.0°C. Analysis calculated: C, 74.41; H, 7.24; N, 10.41. Found: C, 74.38; H, 7.29; N, 10.36.
29y		mp 92.0-93.0°C ESI (m/z) 404

Example 30**N-(3,3-Dimethylbutyl)-6-(hexyloxy)-3-pyridine carboxamide**

A. 6-(Hexyloxy)-3-pyridinecarboxylic acid hexyl ester and 1-Heptyl-1,6-dihydro-6-oxo-3-pyridinecarboxylic acid hexyl ester

A stirred mixture of 6-hydroxynicotinic acid (10.0 g, 0.072 mole) in dimethylformamide (100 mL) under argon at room temperature was treated with 1-bromohexane (30 mL, 35.6 g, 0.216 mole) and powdered potassium carbonate (49.7 g, 0.36 mole) and heated at 110°C for 3 days.

- 15 The reaction mixture, diluted with ethyl acetate, was washed with water and brine, dried (anhydrous magnesium sulfate) and concentrated. Flash chromatography on silica gel and elution with ethyl acetate/hexanes (1:8) gave **2** (4.77 g, 21.5%) and **3** (14.86g, 67.5%).

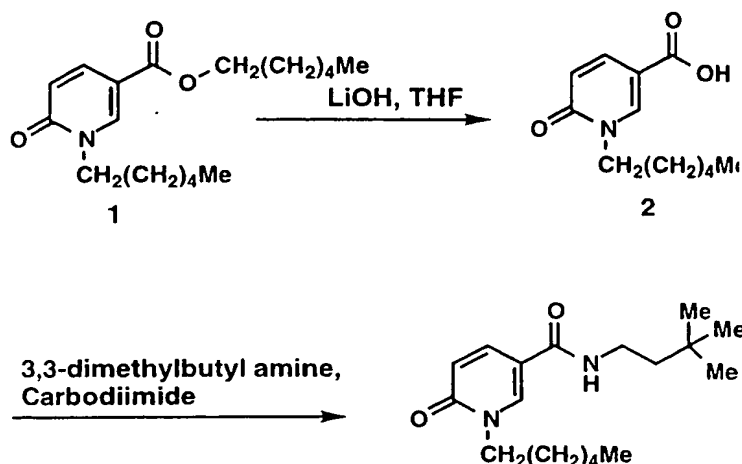
B. 6-(Hexyloxy)-3-pyridinecarboxylic acid

A solution of compound 2 (1.0 g, 3.25 mmole) in tetrahydrofuran (15 mL) was treated with 1 N lithium hydroxide (6.5 mL) and heated at reflux for 24 hours. Solvent was removed *in vacuo* and the residue was partitioned between ethyl acetate and water. The aqueous fraction was acidified with 6 N hydrochloric acid and extracted with ethyl acetate (2x). The combined organic extracts were washed with water and brine, dried (anhydrous magnesium sulfate) and concentrated to give 4 (480 mg, 66%), mp 90-93°C.

C. N-(3,3-Dimethylbutyl)-6-(hexyloxy)-3-pyridine carboxamide

A solution of compound 4 (223 mg, 1.0 mmol) in dimethylformamide (2.5 mL) under argon at room temperature was treated with 3,3-dimethylbutyl amine (111 mg, 1.1 mmol), ethyl-3-(3-dimethylamino)-propylcarbodiimide hydrochloride (327 mg, 1.1 mmol) and hydroxybenzotriazole monohydrate (148 mg, 1.1 mmol) and stirred overnight. The reaction mixture was diluted with ethyl acetate and washed with 10% citric acid, water, dilute sodium bicarbonate, water and brine, dried (anhydrous magnesium sulfate) and concentrated to give an oil. Flash chromatography on silica gel and elution with ethyl acetate/hexanes (1:4) gave an oil (274 mg) that slowly solidified, mp 75-76.5°C.

Analysis calculated for C₁₈H₃₀N₂O₂: C, 70.55; N, 9.87; N, 9.14. Found: C, 70.46; H, 10.06; N, 9.02.

Example 31**N-(3,3-Dimethylbutyl)-1-hexyl-1,6-dihydro-6-oxo-3-pyridinecarboxamide****A. 1-Hexyl-1,6-dihydro-6-oxo-3-pyridinecarboxylic acid**

This compound was prepared from compound 1-Hexyl-1,6-dihydro-6-oxo-3-pyridinecarboxylic acid hexyl ester by the same procedure (lithium hydroxide, THF) as described for the preparation of compound 4 from 2 (Example 30, part B).

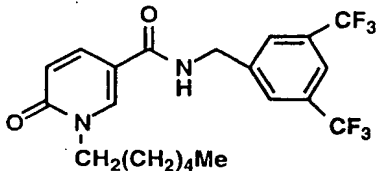
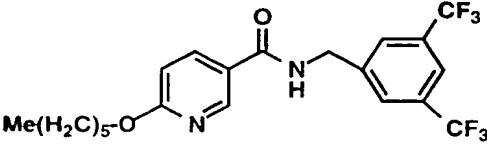
B. N-(3,3-Dimethylbutyl)-1-hexyl-1,6-dihydro-6-oxo-3-pyridinecarboxamide

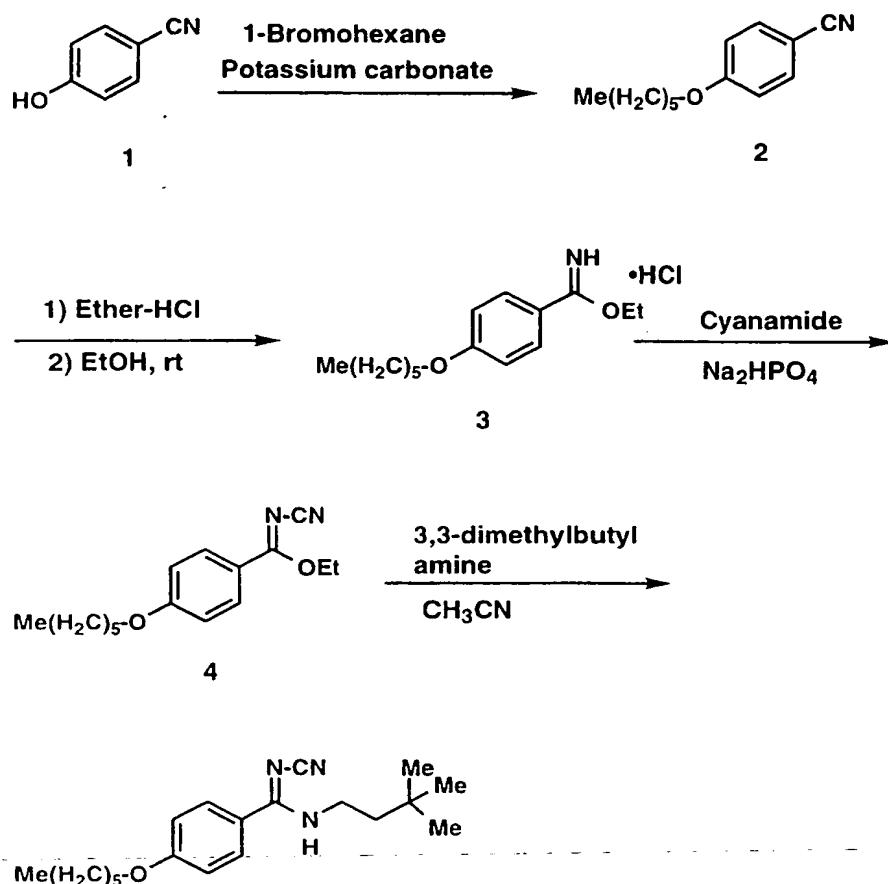
The title compound was prepared from compound 2 by the same procedure as described for the preparation of the title compound of Example 30, part C: mp 80-83°C.

Analysis calculated for $C_{18}H_{30}N_2O_2$: C, 70.55; N, 9.87; N, 9.14. Found:

C, 70.41; H, 9.91; N, 9.37.

Using methodology analogous to that described for the title compounds of Examples 30 and 31, the compounds of Examples 32 and 33 were prepared.

Example #	Structure	Characterization
32	 <chem>CCCCCN1C(=O)C=CC=C1C(=O)NCC2=CC=C(C(F)(F)F)C(F)(F)F2</chem>	$C_{21}H_{22}F_6N_2O_2$: m/e = 448. Analysis calculated: C, 56.25; H, 4.95; N, 6.25; F, 25.42. Found: C, 56.21; H, 4.80; N, 6.16; F, 25.72.
33	 <chem>CCCCCOc1cc(C(=O)NCC2=CC=C(C(F)(F)F)C(F)(F)F2)ccn1</chem>	$C_{21}H_{22}F_6N_2O_2 \cdot 0.25$ hexane: mp 92-95°C. Analysis calculated: C, 57.46; H, 5.47; N, 5.96; F, 24.24. Found: C, 57.46; H, 5.21; N, 6.01; F, 24.04.

Example 34**N-Cyano-N'-(3,3-dimethylbutyl)-4-(hexyloxy)benzene-carboximidamide****A. 4-(Hexyloxy)benzonitrile**

A solution of 4-cyanophenol (20 g, 168 mmol) in dimethylformamide was treated with potassium carbonate (2 eq., 338 mmol, 47 g) and 1-bromohexane (1.1 eq., 185 mmol, 35 mL) and stirred at room temperature for 18 hours. The solid was filtered off and the solution was partitioned between ethyl acetate and 1 N HCl. The organic phase was washed with brine, dried over MgSO_4 , filtered and the solvent was removed to give compound 2 (45 g, >100%).

B. 4-(Hexyloxy)benzenecarboximidic acid ethyl ester

A solution of compound 2 (5.00 g, 24.6 mmol) in diethyl ether was cooled to 0°C and treated slowly with excess freshly prepared cold saturated ethereal-hydrochloric acid (g). The reaction mixture was
5 treated with ethanol (30 mL) and allowed to warm to room temperature and stirred for 18 hours. The solvent was removed to give compound 3 as an oil (4.09 g, 85%).

C. N-Cyano-4-(hexyloxy)benzenecarboximidic acid ethyl ester

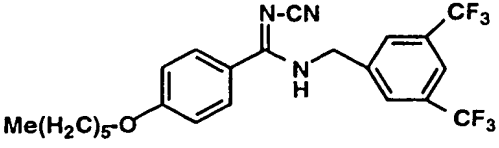
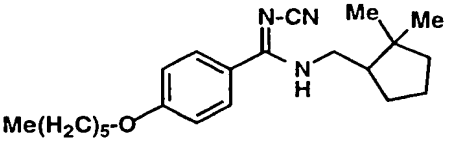
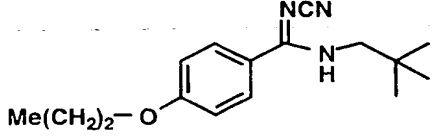
10 A solution of compound 3 (1.0 g, 4.0 mmol) in water/THF (10 mL, 1:1) was treated with sodium dihydrogenphosphate (1.13 g, 8 mmol, in 10 mL water) followed by an aqueous solution of cyanamide (202 mg, 4.8 mmol in 2 mL water). The biphasic solution was stirred vigorously for
15 24 hours. The solution was partitioned between ethyl acetate and saturated sodium bicarbonate solution. The organic phase was washed with brine, dried over magnesium sulfate and the solvent was removed to give compound 4 as a white solid (1.01g, 87%).

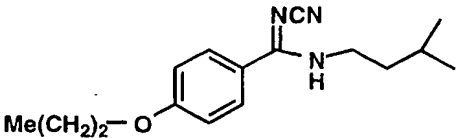
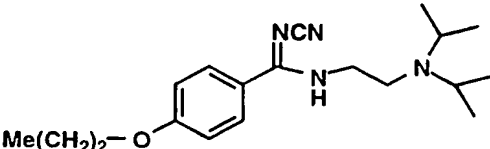
**D. N-Cyano-N'-(3,3-dimethylbutyl)-6-(hexyloxy)-
20 benzenecarboximidamide**

A solution of compound 4 (200 mg, 0.73 mmol) in acetonitrile was treated with 3,3-dimethylbutyl amine (118 µl, 0.875 mmol) and heated to 50°C in a sand bath for 18 hours. The solution was partitioned between ethyl acetate and 10% citric acid solution. The organic phase was
25 washed with brine, dried over magnesium sulfate and evaporated. The residue was purified by flash column chromatography on silica gel using 20% ethyl acetate in hexane to give the title product (90 mg, 38%), mp 89-90°C. Analysis calculated for C₂₀H₃₁N₃O: C, 72.91, H, 9.48, N, 12.75. Found: C, 72.77, H, 9.47, N, 12.66.

30

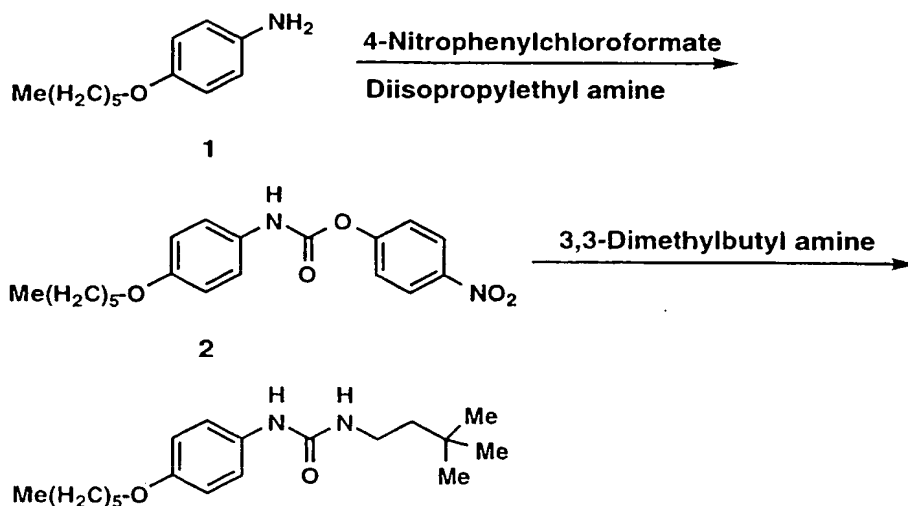
Using methodology analogous to that described for the title compound of Example 34, the compounds of Examples 35 and 36 were prepared.

Example #	Structure	Characterization
35		$C_{23}H_{23}F_6N_3O$: mp 118-129°C. Analysis calculated: C, 58.60; H, 4.92; N, 8.91; F, 24.18. Found: C, 58.99; H, 4.67; N, 8.76; F, 23.93.
36		$C_{22}H_{33}N_3O \cdot 0.03EtO$ Ac. Analysis calculated: C, 73.18; H, 9.67; N, 12.12. Found: C, 73.69; H, 9.67; N, 11.62.
36a		$C_{19}H_{29}N_3O$: mp 69- 71°C. Analysis calculated: C, 72.34; H, 9.27; N, 13.31. Found: C, 72.11; H, 9.36; N, 13.21.

36b		$C_{19}H_{29}N_3O$: mp 106-107°C. Analysis calculated: C, 72.34; H, 9.27; N, 13.31. Found: C, 72.08; H, 9.35; N, 13.19.
36c		$C_{22}H_{36}N_4O \cdot 0.17H_2O$: mp 106-107°C. Analysis calculated: C, 70.93; H, 9.75; N, 14.92. Found: C, 70.35; H, 9.75; N, 14.44.

Example 37

N-(3,3-Dimethylbutyl)-N'-[4-(hexyloxy)phenyl]urea



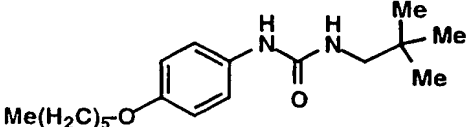
A. [4-(Hexyloxy)phenyl]carbamic acid 4-nitrophenyl ester

A solution of 4-hexyloxyaniline (1.00 g, 5.17 mmol) in dichloromethane (10 mL) was treated with diisopropyl-ethylamine (1.80 mL, 10.3 mmol) followed by p-nitrophenylchloroformate (1.05 g, 5.71 mmol). The reaction mixture was stirred for 3 hours at room temperature and diluted with ethyl acetate. The ethyl acetate solution was washed with hydrochloric acid and brine. After drying over magnesium sulfate the solvent was removed to yield a brown oil. Purification by flash chromatography on silica gel eluted with 15% acetone in hexanes provided compound 2 as a tan solid (241 mg, 13%).

B. N-(3,3-Dimethylbutyl)-N'-[4-(hexyloxy)phenyl]urea

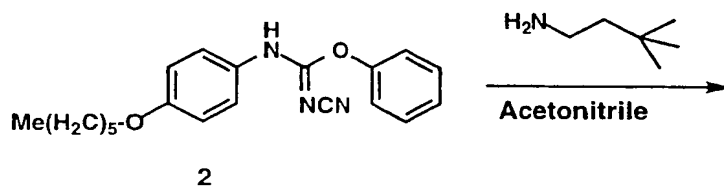
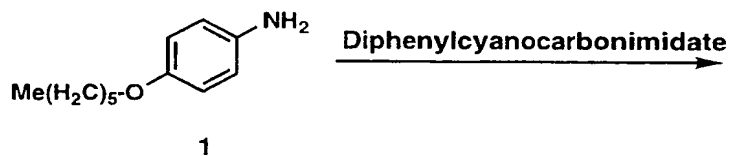
To a solution of compound 2 (90 mg, 0.25 mmol) in tetrahydrofuran (1 mL) was added 3,3-dimethylbutylamine (41 μ L, 0.30 mmol). The reaction was stirred for 18 hours and then purified by flash chromatography on silica gel eluted with 5% 2-propanol in hexanes to provide the title compound as a white solid (75 mg, 94%): mp 103-104°C. Analysis calculated for $C_{19}H_{32}N_2O_2 \cdot 0.14 H_2O$: C, 70.65; H, 10.07; N, 8.67. Found: C, 70.67; H, 10.12; N, 8.52.

Using methodology analogous to that described for the title compound of Example 37, the compound of Example 38 was prepared.

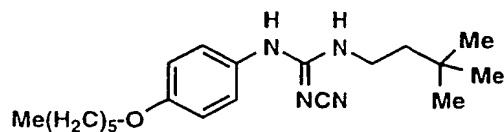
Example #	Structure	Characterization
38		$C_{18}H_{30}N_2O_2 \cdot 0.17$ H_2O : mp 108-109°C. Analysis calculated: C, 69.85; H, 9.88; N, 9.05. Found: C, 69.84; H, 9.85; N, 8.89.

Example 39 (BMS-207307)**N-Cyano-N'-(3,3-dimethylbutyl)-N'-[4-(hexyloxy)phenyl]-guanidine**

5



10

**A. N-Cyano-N'-[4-(hexyloxy)phenyl]carbamimidic acid phenyl ester**

A solution of 4-hexyloxyaniline (1.0 g, 5.17 mmol) in tetrahydrofuran (2 mL) was added to a -25°C solution of potassium
 15 hydride (652 mg of 35% in oil, 5.6 mmol, washed 3 times with hexane) in

dry tetrahydrofuran (2 mL) and was stirred for 30 minutes. The reaction mixture was then treated with diphenylcyanocarbonimide (1.48 g, 6.2 mmol) and stirred overnight while warming to room temperature. The reaction mixture was quenched with water and then partitioned between ethyl acetate and 10% citric acid solution. The organic phase was washed with brine, dried over MgSO_4 , filtered and the solvent was removed. The residue was purified first by flash column chromatography on silica gel using 25% ethyl acetate/hexane as the mobile phase and the product was recrystallized from isopropyl ether/hexane/ CH_2Cl_2 to give compound 2 as white needles (424 mg, 24%).

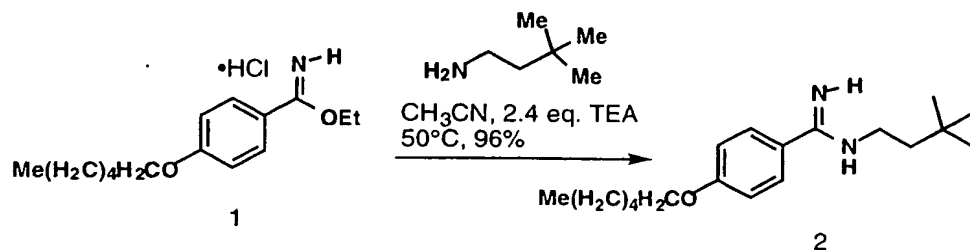
B. N-Cyano-N'-(3,3-dimethylbutyl)-N''-[4-(hexyloxy)phenyl]-guanidine

A solution of compound 2 (400 mg, 1.19 mmol) in acetonitrile/isopropanol (2 mL of 1:1 mixture) was treated with 3,3-dimethylbutylamine (240 μL , 1.79 mmol) and heated to 60°C in a sand bath for 24 hours. The solution was cooled to room temperature, diluted with ethyl ether and washed with 1N HCl. The organic phase was washed with brine, dried over MgSO_4 and the solvent was removed. The residue was crystallized from EtOAc to give the title compound as a colorless solid (168 mg, 41%), mp 183-184°C. Analysis calculated for $\text{C}_{20}\text{H}_{32}\text{N}_4\text{O}$: C, 69.93; H, 9.36; N, 16.26. Found: C, 69.54; H, 9.47; N, 16.05.

Using methodology analogous to that described for the title compound of Example 39, the compounds of Examples 40 to 42 were prepared.

Example #	Structure	Characterization
40	<chem>CCCCCOc1ccc(NC(=N)NCC(C)(C)C)cc1</chem>	$C_{19}H_{30}N_4O$: mp 119-120°C. Analysis calculated: C, 69.06; H, 9.15; N, 16.95. Found: C, 69.02; H, 9.16; N, 16.88.
41	<chem>CCCCCOc1ccc(NC(=N)NCCN(C)C)cc1</chem>	$C_{22}H_{37}N_5O$: mp 88-89°C. Analysis calculated: C, 68.18; H, 9.61; N, 18.07. Found: C, 68.29; H, 9.48; N, 17.82.
42	<chem>CCCCCOc1ccc(NC(=N)NCCC(C)C)cc1</chem>	$C_{19}H_{30}N_4O$: m/e = 330.
42a	<chem>COCCOc1ccc(NC(=N)NCCc2cc(C(F)(F)F)cc(C(F)(F)F)c2)cc1</chem>	$C_{23}H_{24}N_4F_6O \cdot 0$: mp 122-123 °C. Analysis calculated: C, 56.79; H, 4.97; N, 11.52; F, 23.43. Found: C, 56.79; H, 4.84; N, 11.52; F, 23.10.

Example 43

N-(3,3-Dimethylbutyl)-4-(hexyloxy)benzenecarboximidamide,
monohydrochloride

A solution of the title B compound of Example 34 (500 mg, 1.75 mmol) in acetonitrile was treated triethylamine (2.4 eq., 596 μ l), followed by 3,3-dimethylbutylamine (1.5 eq., 2.6 mmol, 353 μ l) and heated at 50°C for 18 hours. The reaction mixture was partitioned between ethyl acetate and saturated sodium bicarbonate. The organic phase was washed with
 10 brine, dried over anhydrous magnesium sulfate, filtered and concentrated. The residue was purified by recrystallization from isopropyl ether/hexane to give the title compound (510 mg, 96%), mp 235-237°C. Analysis calculated for $C_{19}H_{32}N_2O \cdot 1.22 HCl \cdot 1.12 H_2O$: C, 61.82; H, 9.68; N, 7.59; Cl, 11.72. Found: C, 61.83; H, 9.60; N, 7.54; Cl, 12.14.
 15

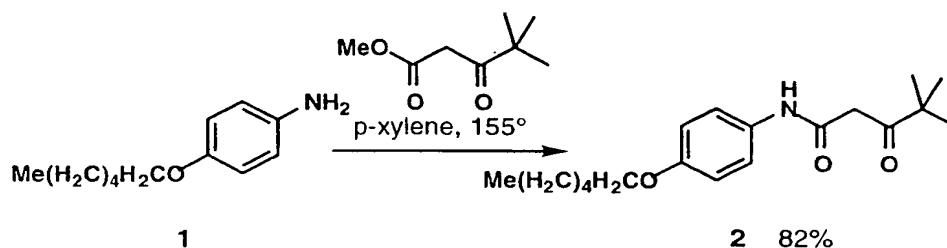
Using the above methodology, the following compounds were prepared.

Example #	Structure	Characterization
44		$C_{22}H_{24}F_6N_2O \cdot 1.0$ HCl: mp 168-169.5°C. Analysis calculated: C, 54.72; H, 5.22; N, 5.80; F, 23.61; Cl, 7.34. Found: C, 55.27; H, 5.34; N, 5.65; F, 23.37; Cl, 7.41.

Example 45

N-[4-(Hexyloxy)phenyl]-4,4-dimethyl-3-oxopentanamide

5

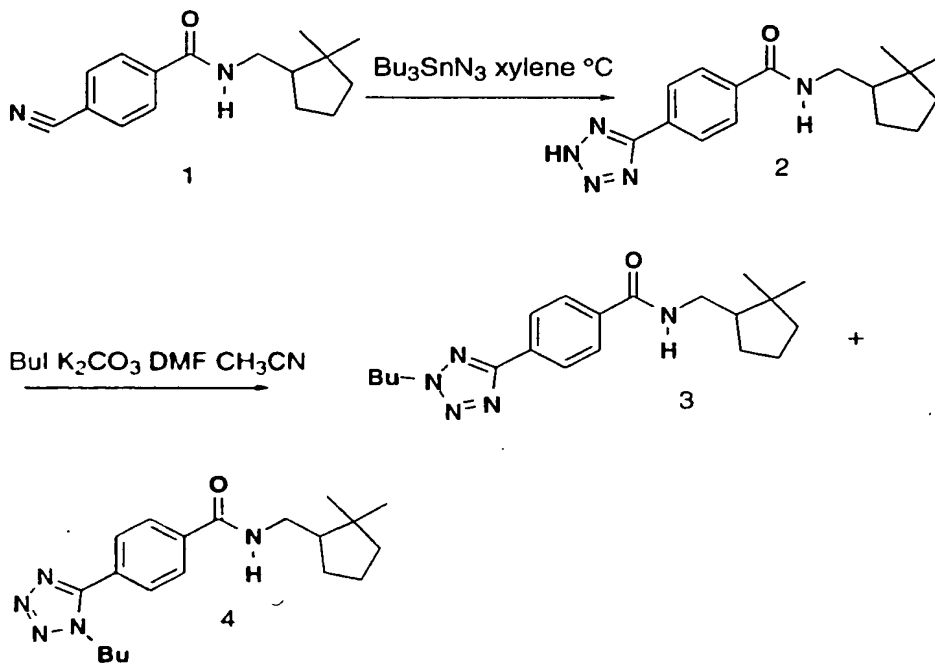


A. N-[4-(Hexyloxy)phenyl]-4,4-dimethyl-3-oxopentanamide

4-Hexyloxyaniline (1, 1.00 g, 5.17 mmol) and methyl-4,4-dimethyl-3-oxopentanoate (1.65 mL, 10.3 mmol) were suspended in 5 mL p-xylene and heated to reflux for 17 hours. The reaction was cooled to room temperature and purified by flash chromatography on silica gel eluting with 10% acetone, hexane to yield a brown solid. Recrystallization from hexane provided 1.35 g (82%) of a white solid. mp 70-72°C; Analysis calculated for $C_{19}H_{29}NO_3$: C, 71.44; H, 9.15; N, 4.38. Found: C, 71.72; H, 9.42; N, 4.30.

Example 46

4-(2-Butyl-2H-tetrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide and 4-(1-butyl-1H-tetrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)methyl]benzamide



A. 4-(1H-Tetrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide

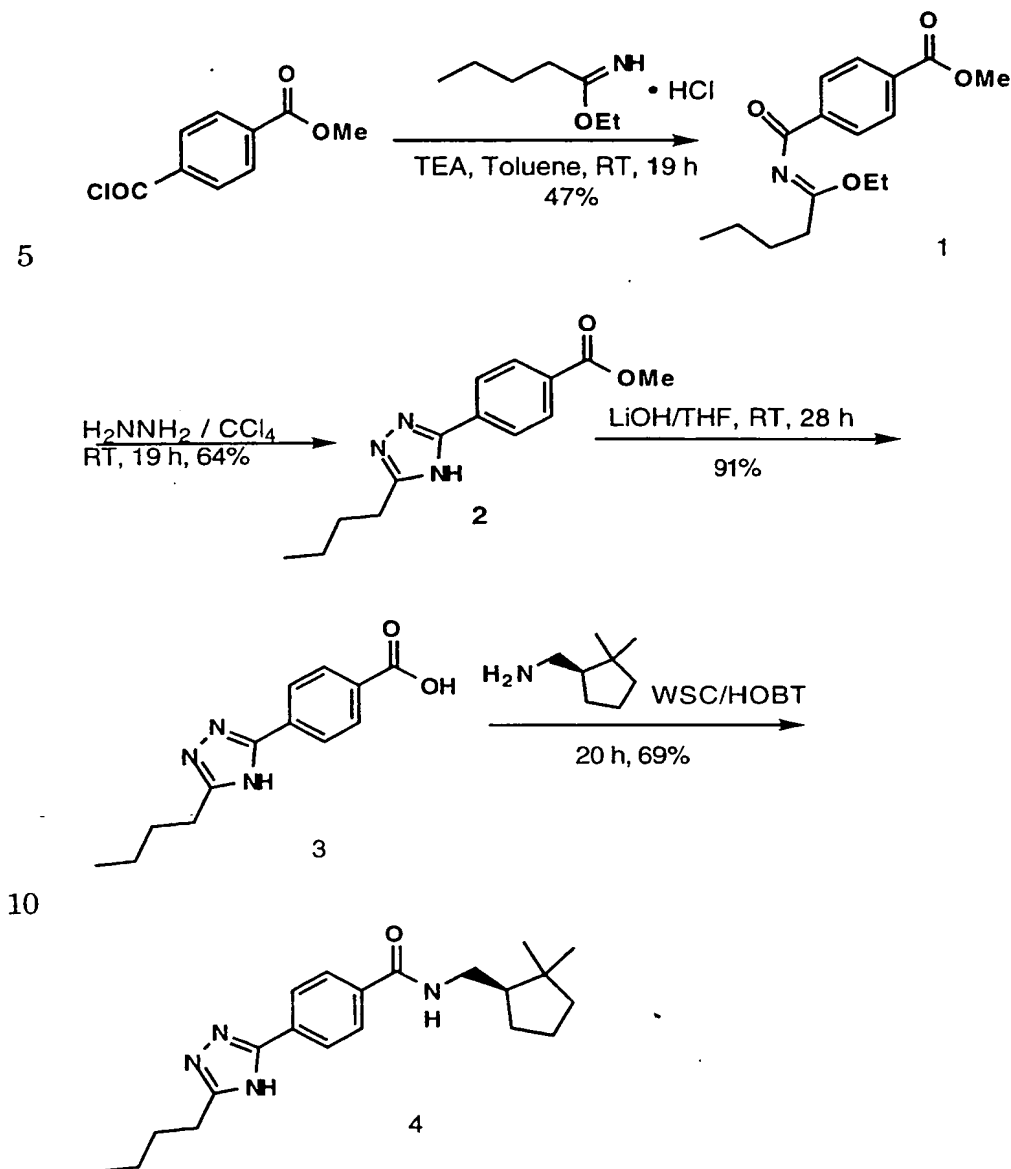
Azidotributyl stannane (5.2 g, 4.4 mL, 15.8 mmol) was added to a solution of the title compound of Example 29b (0.60 g, 2.3 mmol) in xylene (5.1 mL) and the mixture was stirred at 120°C for 15 hours. After cooling, the reaction was diluted with CH_2Cl_2 (40 mL) and 1N HCl (10 mL) and stirred vigorously for 30 minutes. The solid was filtered and rinsed with alternating portions of CH_2Cl_2 (3 x 20 mL) and 1N HCl (3 x 20 mL) and dried under high vacuum overnight to afford the title compound as a white solid.

B. 4-(2-Butyl-2H-tetrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)-methyl] benzamide and 4-(1-butyl-1H-tetrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)methyl]benzamide

- 5 A mixture of the title A compound (0.60 g, 2.0 mmol), iodobutane (0.38 g, 0.24 mL, 2.1 mmol), K_2CO_3 (0.40 g, 3.0 mmol), and DMF (5.0 mL) in acetonitrile (20 mL) was stirred at 55°C. After 19 hours, the reaction was transferred to a separatory funnel with EtOAc/1N HCl. The mixture was extracted with EtOAc (2 x 40 mL), washed with brine, dried
- 10 (MgSO₄) and concentrated *in vacuo*. Flash chromatography (silica, 50 mm, 25% EtOAc/hexane) afforded the 4-(2-butyl-2H-tetrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)methyl] benzamide (0.57 g, 81%): mp 121-123°C. Analysis calculated for C₂₀H₂₉N₅O-C, 67.58; H, 8.22; N, 19.7; Found: C, 67.48; H, 8.46; N, 19.68.
- 15 Further elution afforded 4-(1-butyl-1H-tetrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)methyl]benzamide (46 mg, 6%): mp 106-109°C. Analysis calculated for C₂₀H₂₉N₅O•H₂O: C, 65.27; H, 8.32; N, 19.03. Found: C, 65.32; H, 8.31; N, 18.63.

Example 47

(R)-N-[(2,2-Dimethylcyclopentyl)methyl]-4-(5-butyl-1,2,4H-triazol-3-yl)benzamide



A. 4-[(1-Ethoxypentylidene)amino]carbonyl]benzoic acid methyl ester

To a solution of ethyl pentanimidate hydrochloride (9 g, 54 mmol) and triethyl amine (17 mL, 120 mmol) in toluene (110 mL) at 0°C was added, over 20 minutes, a solution of terephthalic acid, monomethyl ester chloride (10.8 g, 54 mmol) in toluene (21 mL). After stirring at ambient temperature for 19 hours, the reaction was filtered and the solids were rinsed with toluene (200 mL). The combined filtrates were concentrated *in vacuo*. Flash chromatography (silica gel, CH₂Cl₂) of a portion of this material afforded the title compound (1.9 g, 47%).

B. 4-(5-Butyl-1H-1,2,4-triazol-3-yl)benzoic acid methyl ester

Hydrazine (0.23 g, 0.22 mL, 7.0 mmol) was added to a solution of the title A compound (1.9 g, 6.4 mmol) in CCl₄ (32 mL). After standing at ambient temperature for 19 hours, the reaction was transferred to a separatory funnel containing CH₂Cl₂/H₂O. The mixture was extracted with CH₂Cl₂ (2 x 50 mL) and 10% MeOH/CH₂Cl₂ (3 x 30 mL). The combined organic layers were dried (MgSO₄) and concentrated *in vacuo*. Flash chromatography (silica gel, 50% EtOAc/hexane) gave the title compound (1.06 g, 64%).

C. 4-(5-Butyl-1,2,4-1H-triazol-3-yl)benzoic acid

Lithium hydroxide (1 N, 8.2 mL, 8.2 mmol) was added to a solution of the title B compound (1.06 g, 4.1 mmol) in THF (45 mL). After stirring at ambient temperature for 28 hours, the reaction was evaporated *in vacuo*. The aqueous residue was acidified with 1 N HCl to pH 3 to 4. The resulting solid was filtered, washed with H₂O, and dried under high vacuum to afford the title compound (0.90 g, 91%).

D. (R)-N-[(2,2-Dimethylcyclopentyl)methyl]-4-(5-butyl-1,2,4-1H-triazol-3-yl)benzamide

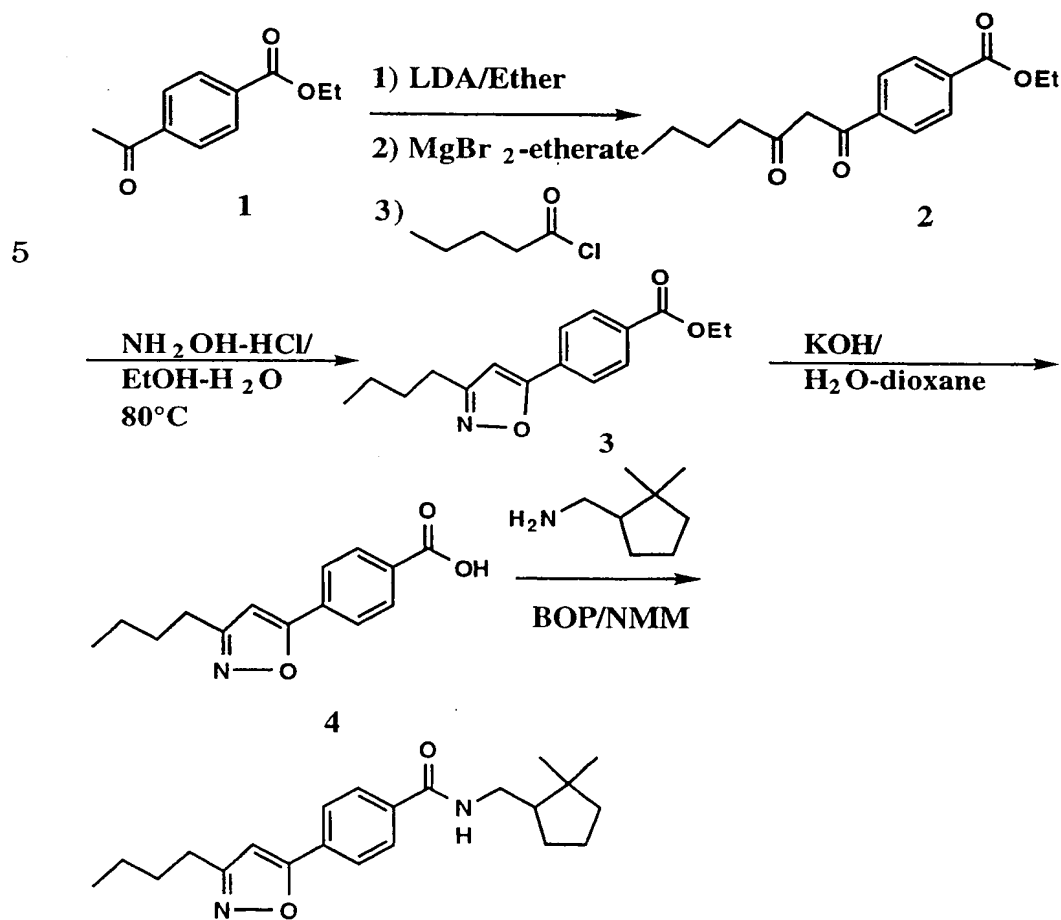
1-Hydroxybenzotriazole hydrate (70 mg, 0.51 mmol) and 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (WSC, 156 mg, 0.51 mmol) were added to a solution of the title C compound (123 mg, 0.50 mmol) in CH_2Cl_2 (2.2 mL) and DMF (0.57 mL) stirring at ambient temperature. After stirring for 30 minutes, (R)-((2,2-dimethylcyclopentyl)methyl)amine, hydrochloride (82 mg, 0.50 mmol) in CH_2Cl_2 (0.64 mL) was added followed by triethylamine (0.070 mL, 0.50 mmol). After stirring for 20 hours, the reaction was transferred to a separatory funnel with ether/ H_2O and the aqueous layer acidified with 1N HCl. The mixture was extracted with ether (2 x 80 mL), washed with saturated NaHCO_3 , brine, dried (MgSO_4) and concentrated *in vacuo*. Flash chromatography (silica gel, 50% EtOAc/ CH_2Cl_2) afforded the title compound (0.12 g, 69%): mp 182.0-184.5°C. Analysis calculated for $\text{C}_{21}\text{H}_{30}\text{N}_4\text{O} \cdot 0.27 \text{ H}_2\text{O}$: C, 70.20; H, 8.57; N, 15.59. Found: C, 70.20; H, 8.80; N, 15.43.

Using the procedure described in Example 47, the following compounds were prepared.

Example #	Structure	Characterization
48		$C_{21}H_{30}N_4O \cdot 0.18$ H_2O : ESI (m/z) 355. Analysis calculated: C, 70.50; H, 8.55; N, 15.66. Found: C, 70.50; H, 8.72; N, 15.44.

Example 49

4-(3-Butyl-5-isoxazolyl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide



A. 4-(3-Butyl-5-isoxazolyl)benzoic acid ethyl ester

A solution of ethyl 4-acetylbenzoate (2.93 g, 15.3 mmol) in 25 mL ether was added to a solution of lithium diisopropylamide (0.805 M in THF-hexane, 20.85 mL, 16.79 mmol) in 200 mL of ether at -70°C. The mixture was stirred at -78°C for 30 minutes, magnesium bromide etherate (3.94 g, 15.26 mmol) was added and the reaction mixture was stirred for 30 minutes followed by the addition of valeryl chloride (1.47 g, 12.2 mmol) in 3 mL ether. Stirring was continued at -78°C for an additional 30 minutes. The mixture was allowed to come to room temperature, quenched by adding saturated ammonium chloride and acidified (pH = 3.0) by adding 20% sulfuric acid. The mixture was extracted with ethyl acetate, the organic layer was dried over magnesium sulfate, concentrated, and the residue subjected to flash chromatography (silica gel/hexane-ethyl acetate 95:5) to afford the title compound (1.35 g).

B. 4-(3-Butyl-5-isoxazolyl)benzoic acid ethyl ester

A solution of the title A compound (150 mg, 0.54 mmol) in 1.5 mL ethanol was treated with a solution of hydroxylamine hydrochloride (41.4 mg, 0.6 mmol) in 0.15 mL of water. The mixture was heated at 80°C in a sealed tube for 2 hours, concentrated, and the residue partitioned between water and methylene chloride. The methylene chloride layer was dried over magnesium sulfate and concentrated to afford the desired product (145 mg).

C. 4-(3-Butyl-5-isoxazolyl)benzoic acid

A solution of the the title B compound (145 mg) in 6 mL dioxane was treated with 10% aqueous potassium hydroxide (6 mL) and the mixture was stirred at room temperature for 14 hours. The mixture was diluted with water, washed with ether, the aqueous layer was acidified with 10% sodium hydrogen sulfate and extracted with ethyl acetate. The ethyl

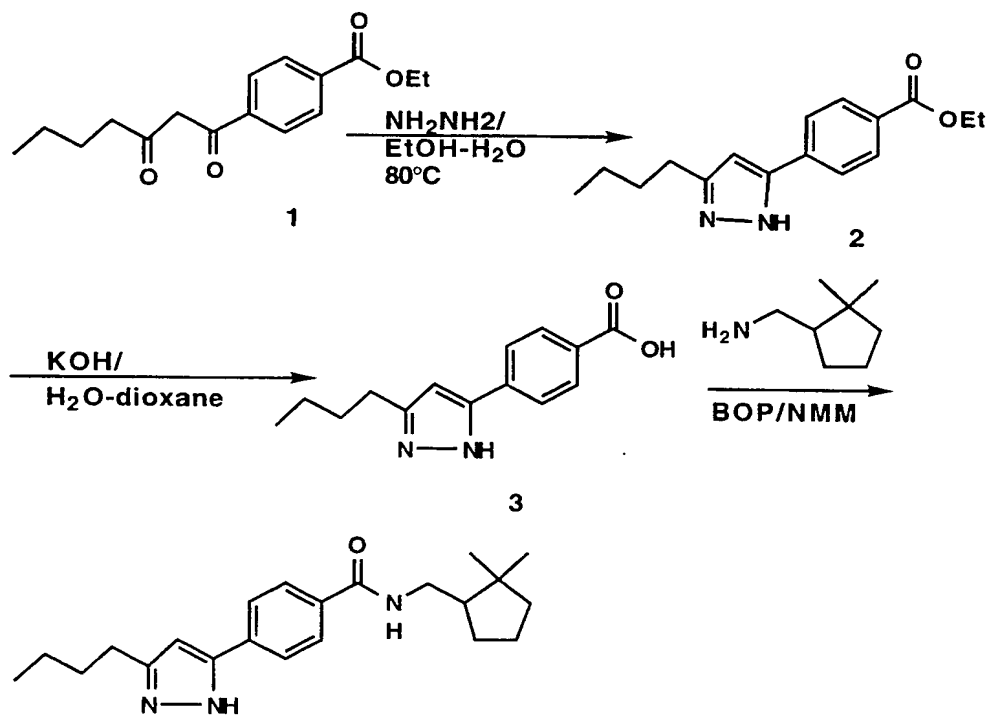
acetate layer was dried over magnesium sulfate, concentrated to give the desired compound (88 mg) as a white solid.

D. 4-(3-Butyl-5-isoxazolyl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide

The title C compound was sequentially treated with benzotriazol-1-yloxytris(dimethylamino)phosphonium hexafluorophosphate (BOP) (174.4 mg, 0.395 mmol), 2,2-dimethylcyclopentylmethyl amine (59 mg, 0.359 mmol; the title B compound of Example 1) and N-methylmorpholine in 3 mL DMF. The mixture was stirred at room temperature for 14 hours, diluted with water, stirred for 5 minutes and the resulting precipitate was isolated by filtration. The crude product was recrystallized from hexane to give the title compound (90 mg) as a white solid.

Example 50

4-(3-Butyl-1H-pyrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide

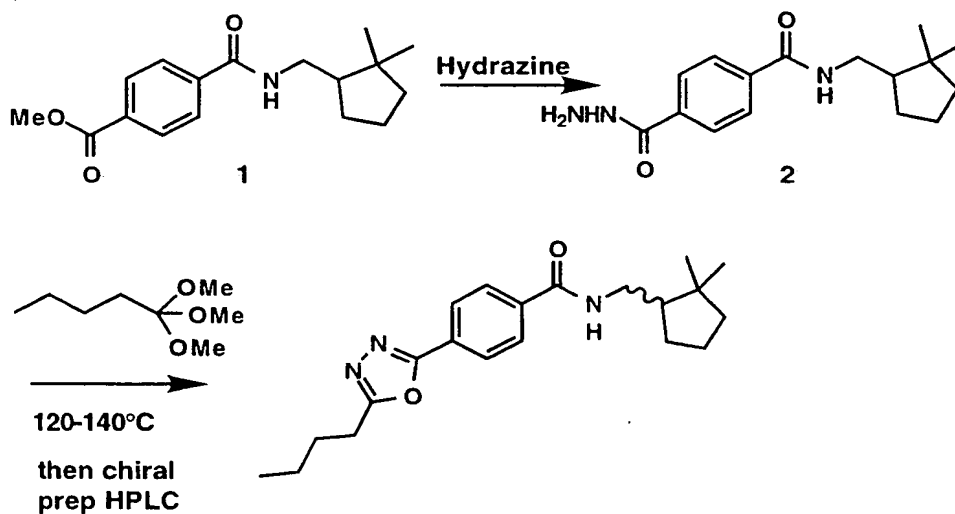


A. 4-(3-Butyl-1H-pyrazol-5-yl)benzoic acid ethyl ester

This title compound was prepared from compound 1 (title A compound of Example 49) and hydrazine by a similar procedure as described for the title B compound of Example 49.

B. 4-(3-Butyl-1H-pyrazol-5-yl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide

The title A compound was converted to the desired product by a sequence of reactions as described for the preparation of the title compound of Example 49 from the title B compound of Example 49. The product was obtained as a white solid, MS: m/e = 353.

Example 51**15 4-(5-Butyl-1,3,4-oxadiazol-2-yl)-N-[(2,2-dimethylcyclopentyl)-methyl]benzamide, enantiomer A and enantiomer B**

A. 4-[[[(2,2-Dimethylcyclopentyl)methyl]amino]-carbonyl]benzoic acid methyl ester

The title compound was prepared from monomethylterephthalate and 2,2-dimethylcyclopentylmethyl amine (title B compound of Example 1) in a manner as described for the title A compound of Example 24.

B. 4-[[[(2,2-Dimethylcyclopentyl)methyl]amino]-carbonyl]benzoic acid hydrazide

The title A compound (275 mg) was treated with hydrazine monohydrate and heated at 120°C for 2 hours. The reaction mixture was diluted with water, saturated with potassium carbonate and extracted with ethyl acetate. The organic layer was dried over magnesium sulfate and concentrated to give the title compound (270 mg) as a white solid which was used in the next reaction without isolation.

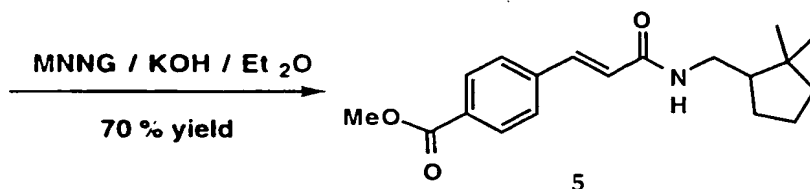
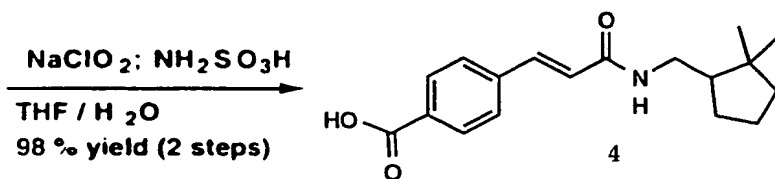
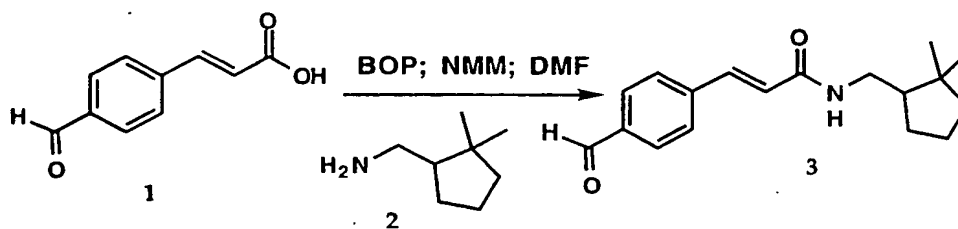
C. 4-(5-Butyl-1,3,4-oxadiazol-2-yl)-N-[(2,2-dimethylcyclopentyl)methyl]benzamide, enantiomer A and enantiomer B

The title B compound was heated in trimethyl orthoacetate (3 mL), at 120°C for 2 hours. The excess trimethyl orthoacetate was removed by purging with a stream of nitrogen at 140°C. The residue was heated at 140°C for 1.5 hours, diluted with methylene chloride, washed with 10% potassium carbonate and the organic layer was dried (magnesium sulfate) and concentrated. The crude product was subjected sequentially to flash chromatography (silica gel/hexane-EtOAc 1:1) and chiral preparative HPLC (Chirapak AD column/hexane-isopropanol-triethylamine 80:20:0.2) to give the two enantiomers: (+)-enantiomer A (104 mg, $[\alpha]_D = +22^\circ\text{C} = 0.36$, methylene chloride, m/e 355) and the (-)-enantiomer B (100 mg, $[\alpha]_D = -19.5^\circ\text{C} = 0.36$, methylene chloride, m/e 355) as white solids.

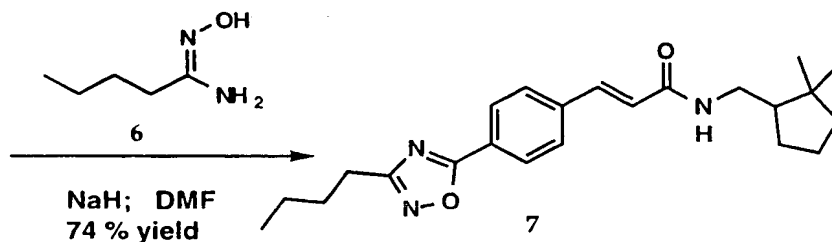
Example 52

3-[4-(3-Butyl-1,2,4-oxadiazol-5-yl)phenyl]-N-[(2,2-dimethylcyclopentyl)methyl]-2-propenamide

5



10



15 A. N-[(2,2-Dimethylcyclopentyl)methyl]-3-(4-formylphenyl)-2-propenamide

To a cold (0°C) solution of 4-formylcinnamic acid (345 mg, 1.96 mmol) and 2,2-dimethylcyclopentylmethylamine, hydrochloride (321 mg, 1.96 mmol ; title B compound of Example 1) in DMF (5 mL) was added

20 benzotriazol-1-yloxytris(dimethylamino)phosphonium

hexafluorophosphate (BOP) (954 mg, 2.16 mmol) and N-methylmorpholine (0.47 mL, 4.31 mmol). The cooling bath was removed and the reaction mixture was stirred at room temperature for 18 hours. The reaction mixture was diluted with dichloromethane and washed
5 with 1N HCl and saturated NaHCO₃ solution. The organic layer was dried (MgSO₄) and concentrated *in vacuo* to yield a light amber gum. MS: (ESI), (M+H)⁺ 286.

B. 4-[3-[[2,2-Dimethylcyclopentyl)methyl]amino]-3-oxo-1-propenyl]benzoic acid
10

To stirred solution of the title A compound in THF/H₂O (15 mL of a 1:1 mixture) at 0°C was added sodium chlorite (329 mg, 3.64 mmol) and sulfamic acid (353 mg, 3.64 mmol). The cooling bath was removed and the reaction was allowed to warm up to room temperature. Once the
15 reaction was complete, the mixture was diluted with dichloromethane and washed with saturated potassium bisulfate solution. The organic layers were separated and the aqueous layer was backwashed with fresh dichloromethane (twice). The combined organic extracts were washed with freshly prepared 2% sodium bisulfite solution, dried (MgSO₄) and
20 concentrated *in vacuo* to afford a light yellow solid (583 mg, 98%), mp 138-140°C.

C. 4-[3-[[2,2-Dimethylcyclopentyl)methyl]amino]-3-oxo-1-propenyl]benzoic acid methyl ester
25

The freshly prepared excess diazomethane was slowly added to a cold (0°C) solution of the title B compound (583 mg, 1.93 mmol) in diethyl ether (50 mL) until a yellow color persisted. The reaction was quenched with a few drops of glacial acetic acid until the yellow color disappeared. The reaction was then concentrated *in vacuo* to afford a yellow gum
30 which was purified by flash chromatography on silica gel (10% EtAc in

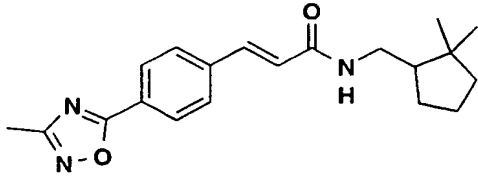
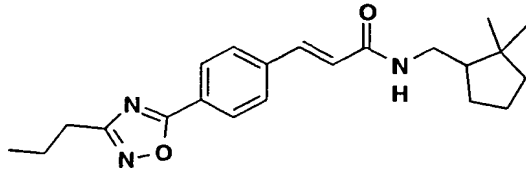
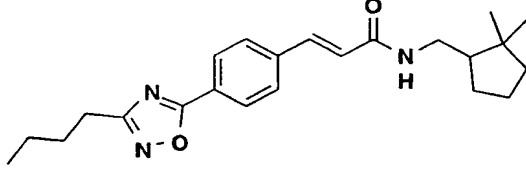
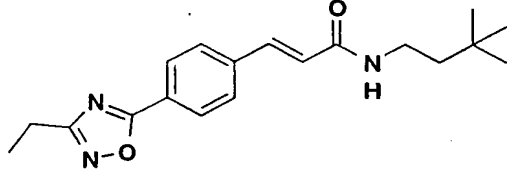
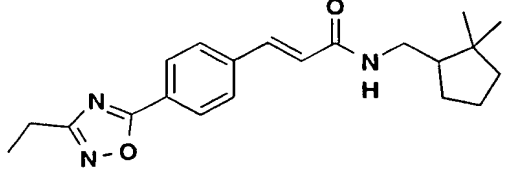
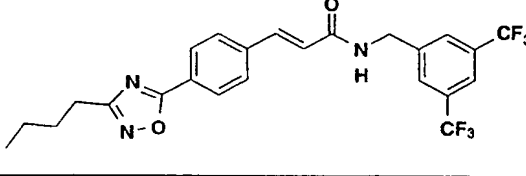
hexane) gave the desired product as a creamy light yellow solid. MS: (ESI) (M+H)⁺ 316.

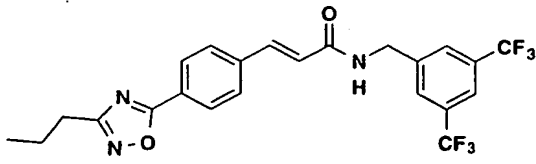
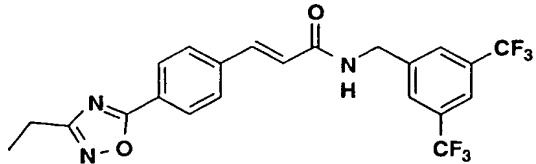
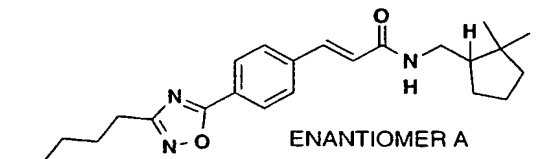
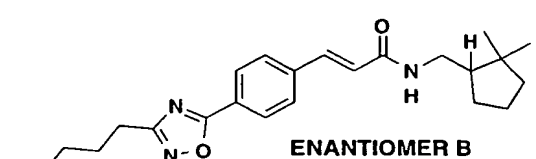
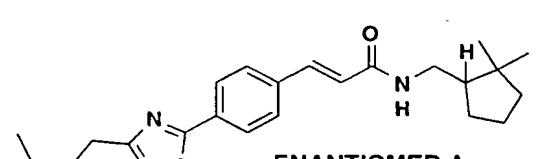
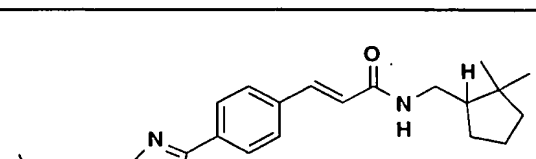
D. 3-[4-(3-Butyl-1,2,4-oxadiazol-5-yl)phenyl]-N-[(2,2-dimethylcyclopentyl)methyl]-2-propenamide

To a solution of the title C compound (100 mg, 0.32 mmol) and n-butylhydroxyamidine (46 mg, 0.40 mmol) in DMF (2 mL) was added NaH (17 mg, 0.70 mmol). After stirring for 18 hours at room temperature, the reaction mixture was diluted with dichloromethane and washed with water. The combined organic extracts were dried (MgSO₄), and concentrated *in vacuo* to yield a yellow solid (90 mg, 74%), mp 144-145°C. MS (ESI): (M+H)⁺ 382.

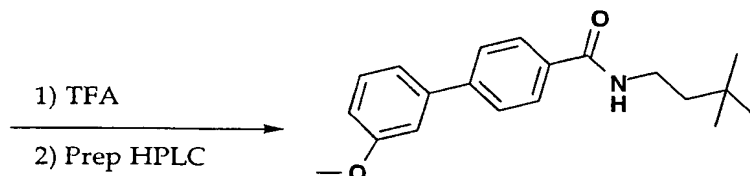
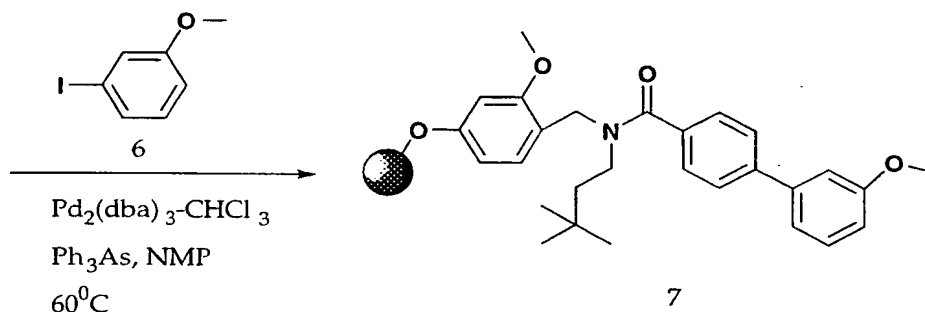
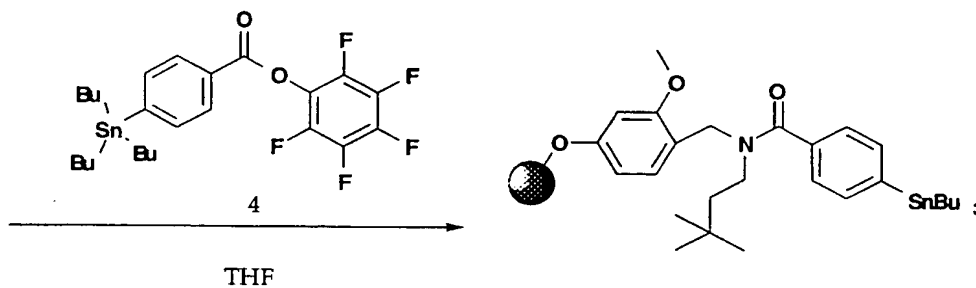
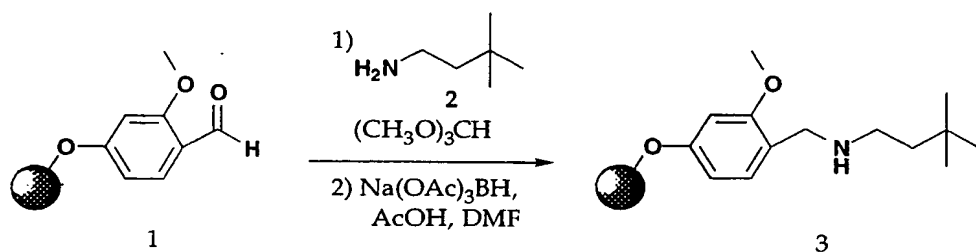
Using the procedures described above the following compounds were prepared.

Example #	Structure	Characterization
53		C ₁₈ H ₂₃ N ₃ O ₂ : mp 185-186°C
54		C ₂₀ H ₂₇ N ₃ O ₂ : mp 162-164°C
55		C ₂₁ H ₂₉ N ₃ O ₂ : mp 164-166°C

Example #	Structure	Characterization
56		$C_{20}H_{25}N_3O_2$: mp 160-161°C
57		$C_{22}H_{29}N_3O_2$: mp 147-149°C
58		$C_{23}H_{31}N_3O_2$: mp 146-148°C
59		$C_{19}H_{25}N_3O_2$: mp 162-163°C
60		$C_{21}H_{27}N_3O_2$: mp 145-146°C
61		$C_{24}H_{21}F_6N_3O_2$: mp 178-179°C

Example #	Structure	Characterization
62		$C_{23}H_{19}F_6N_3O_2$: mp 178-179°C
63		$C_{22}H_{17}F_6N_3O_2$: MS: $M+H = 468$
64	 <p>ENANTIOMER A</p>	$C_{24}H_{33}N_3O_2$: mp 131-132°C
65	 <p>ENANTIOMER B</p>	$C_{24}H_{33}N_3O_2$: mp 130-131°C
66	 <p>ENANTIOMER A</p>	$C_{24}H_{33}N_3O_2$: mp 144-145°C
67	 <p>ENANTIOMER B</p>	$C_{24}H_{33}N_3O_2$: mp 140-141°C

Example 68



A. Resin 1

To a stirred suspension of NaH (5.25 g, 131 mmol) in 100 mL of dry
 15 DMF at 4°C was added a solution of 4-hydroxy-2-methoxybenzaldehyde
 (20.0g, 131 mmol) in 100 mL of dry DMF drop-wise via addition funnel

over a period of 30 minutes. The addition funnel was washed with 30 mL of dry DMF which was added to the reaction flask. After stirring at 4°C for one hour, the ice-water bath was removed and the reaction was allowed to warm to room temperature over the period of 1/2 hour. At the end of this period, tetra-n-butylammonium iodide (6.5 g, 17.6 mmol) was added followed by Merrifield resin (52.8 g, loading = 1.24 mmol/g, 65.5 mmol). The reaction flask was immersed in an oil bath which was heated to 70°C. After 20 hours, the heating bath was removed and the reaction was allowed to cool to room temperature. The reaction was filtered with suction and the residual resin rinsed with water:DMF (3 x 200 mL), DMF (3 x 200 mL), THF (3 x 200 mL), methanol (2 x 200 mL). The resin was dried under (20 mm Hg) for 18 hours to yield 62.86 g of resin 1. Elemental analysis indicated less than 0.1% residual Cl.

B. Resin 2

Resin 1 (10g, loading = 1.08 mmol/g, 10.8 mmol) was weighed into a glass solid phase organic synthesis (SPOS) reaction vessel. The resin was swelled with dry DMF (50 mL) which was drained after 10 minutes. Dry DMF (60 mL) was then added followed by 3,3-dimethylbutylamine (3.0 g, 29.6 mmol). The reaction vessel was agitated on a wrist-action shaker for 10 minutes and trimethyl orthoformate (30 mL) was added. After agitating for 14 hours, the reaction was drained and 50 mL of dry DMF was added. The reaction was agitated for approximately 1 minute and drained. Dry DMF (10 mL) was added followed by sodium triacetoxyborohydride (6.35 g, 30 mmol) and acetic acid (0.60 mL, 1 mmol). After 6 hours of agitation, the reaction was drained and rinsed with DMF (3 x 50 mL), DMF-water (3 x 50 mL), DMF (3 x 50 mL), dichloromethane (3 x 50 mL), methanol (3 x 50 mL) and THF (3 x 50 mL). The resin was used in the next step without characterization.

C. Compound 4

To a stirred solution of pentafluorophenol (7.6 g, 41.3 mmol) in dry dichloromethane at 4°C was added N,N-diisopropylethylamine (19.6 mL, 112.5 mmol) drop-wise via syringe over a period of 5 minutes. After 20 minutes, 4-iodobenzoyl chloride (10.0 g, 37.5 mmol) was added as a solid. The reaction was stirred at 4°C for 2 hours and the ice-water bath was removed. The reaction was allowed to warm to room temperature over a half hour period. TLC analysis indicated complete consumption of starting material. The reaction was poured into 500 mL of dichloromethane which was washed with 1N aq. HCl (3 x 200 mL), water (2 x 200 mL), sat. aq. sodium bicarbonate (3 x 200 mL) and brine (200 mL), dried over magnesium sulfate, filtered and stripped. The product was dried under vacuum to yield 15.35 g (37.0 mmol) of a white waxy solid. This material was dissolved in 100 mL of dry toluene and bis(tributyltin) (28 mL, 55.4 mmol) was added. The reaction mixture was degassed by bubbling with dry nitrogen for a period of 15 minutes. At the end of this period, tetrakis(triphenylphosphine) palladium (0) (427 mg, 0.37 mmol) was added. The reaction flask was fitted with a reflux condenser and immersed in an oil bath which was heated to 120°C. After stirring for 14 hours, the reaction was allowed to cool and filtered through a pad of silica. The solvent was removed *in vacuo*. The residual oil was purified by flash reverse-phase chromatography (YMC-gel, dichloromethane-acetonitrile). The tubes containing the product were pooled and concentrated to yield 16.31g (28.3 mmol, 75.5 % from 4-iodobenzoyl chloride) of 4.

D. Resin 5

Resin 3 (4.3 g, loading = 0.99 mmol/g, 4.3 mmol) was transferred into a glass SPOS reaction vessel. The resin was swelled with 20 mL of dry THF. After 10 minutes the solvent was drained away and 5 mL of

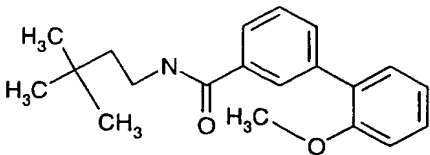
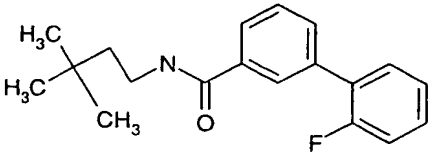
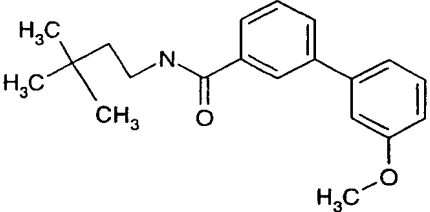
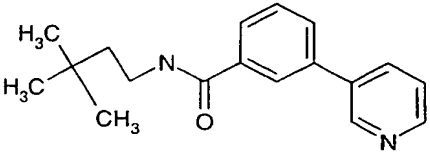
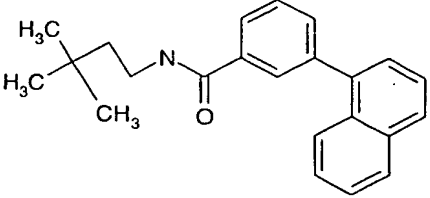
dry THF was added. A solution of compound 4 (4.9 g, 8.5 mmol) in 15 mL of dry THF was added via syringe. The reaction was agitated on a wrist-action shaker for 16 hours. The solvent was drained from the resin which was then washed with THF (3 x 40 mL), dichloromethane (3 x 40 mL), methanol (3 x 40 mL) and THF (3 x 40 mL). The resin was dried *in vacuo* at 20 mm Hg for 18 hr. The resin was used in the next step without further characterization.

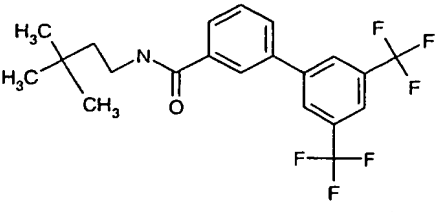
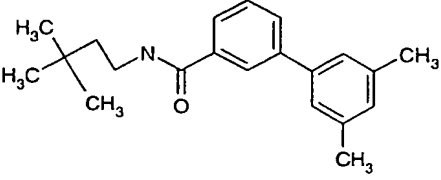
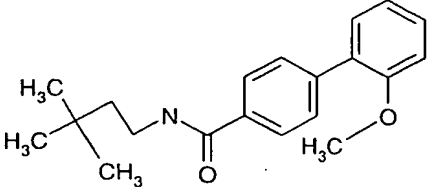
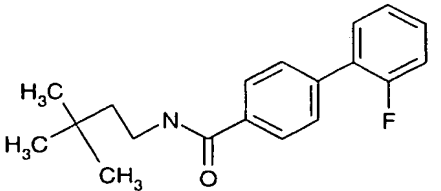
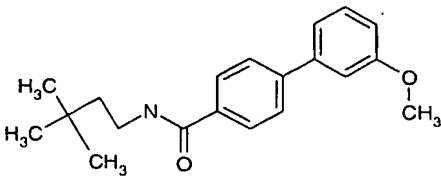
E. Resin 7

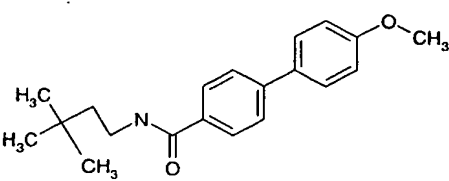
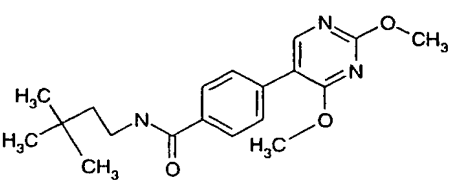
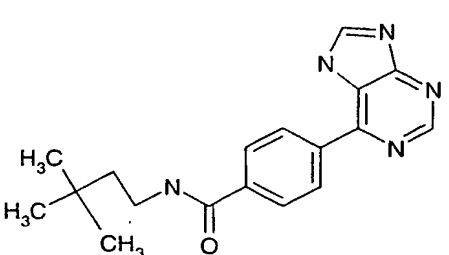
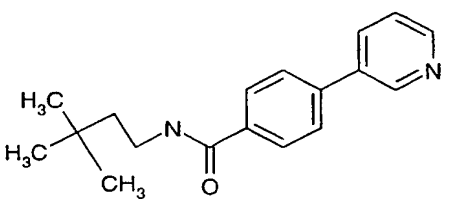
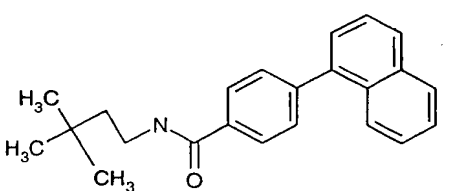
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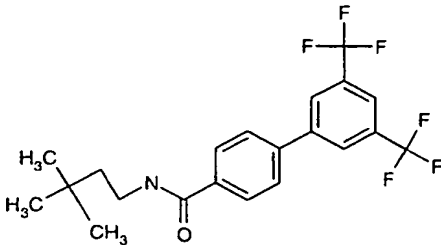
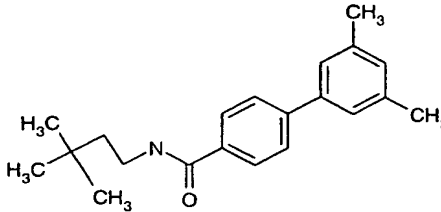
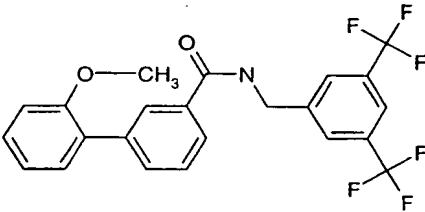
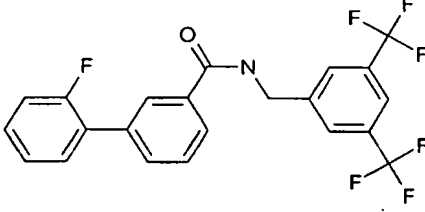
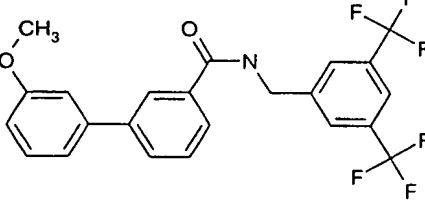
Resin 5 (300 mg, 0.213 mmol) was transferred into a polypropylene reaction tube. Dry NMP (2 mL) was added followed by 3-iodoanisole (169 mg, 0.72 mmol), triphenylarsine (45 mg, 0.15 mmol) and tris(dibenzylideneacetone)dipalladium-(0)-chloroform adduct (35 mg, 0.034 mmol). The reaction was sealed and agitated at 275 rpm on an orbital shaker. The reaction was heated to 55°C over a one hour period. After 14 hours, the reaction mixture was allowed to cool to room temperature. The solvent was drained and the resin was washed with DMF (3 x 5 mL), DMF-water (3 x 5 mL), dichloromethane (3 x 5 mL), THF (3 x 5 mL), methanol (3 x 5 mL), and dichloromethane (3 x 5 mL). The resin was suspended in dichloromethane (0.5 mL) and trifluoroacetic acid (3 mL). After 1 hour, the product was collected into a tared test-tube. The solvent was removed *in vacuo*. The crude product was purified by preparative HPLC using a YMC S3 ODS 20x100 mm column with a 30 - 100% B gradient over 10 minutes at a flow rate of 25 mL/minute (Solvent A: 90% water/ 10% methanol with 0.1% TFA; Solvent B 10% water/ 90% methanol with 0.1% TFA) to provide the title compound (3.7 mg).

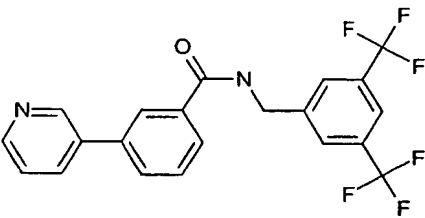
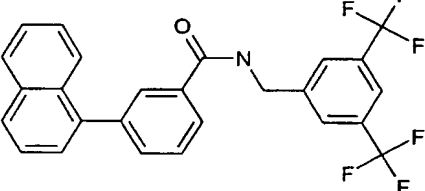
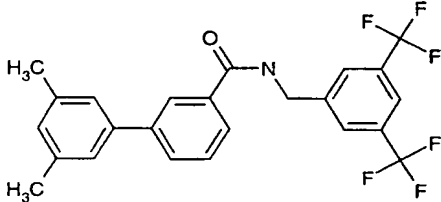
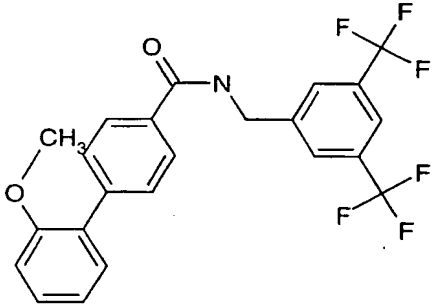
Using the procedure described in Example 68, the following compounds were prepared.

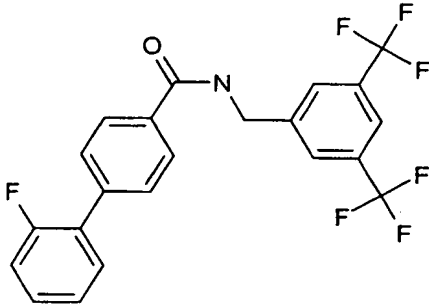
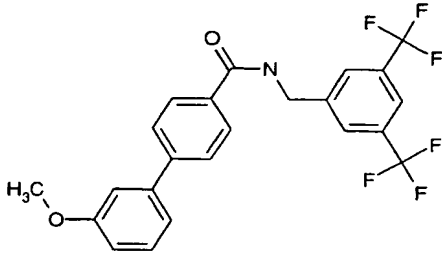
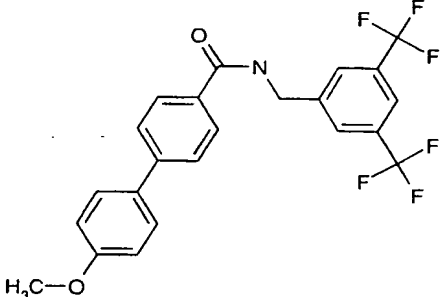
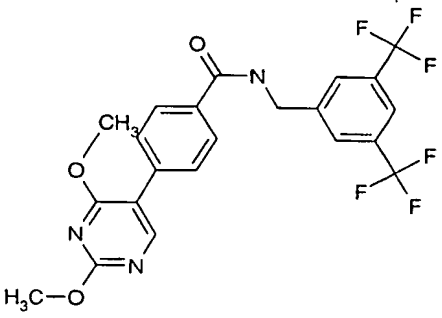
Example #	Structure	Characterization
69		$C_{20}H_{25}NO_2$ m/z 312 (M+H)
70		$C_{19}H_{22}FNO$ m/z 300 (M+H)
71		$C_{20}H_{25}NO_2$ m/z 312 (M+H)
72		$C_{18}H_{22}N_2O$ m/z 283 (M+H)
73		$C_{23}H_{25}NO$ m/z 332 (M+H)

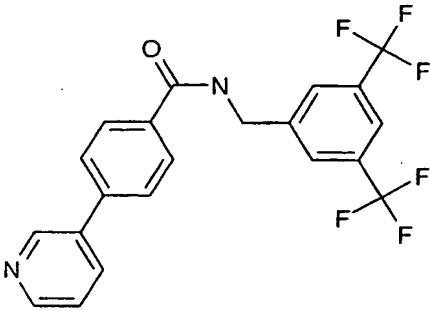
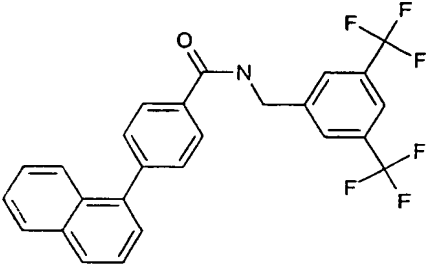
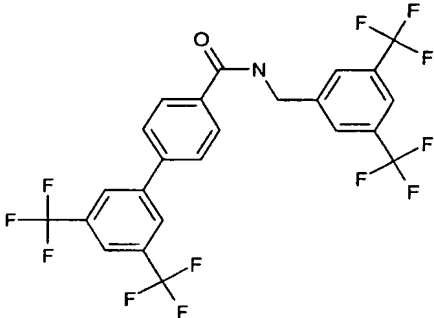
Example #	Structure	Characterization
74		$C_{21}H_{21}F_6NO$ m/z 418 (M+H)
75		$C_{21}H_{27}NO$ m/z 310 (M+H)
76		$C_{20}H_{25}NO_2$ m/z 312 (M+H)
76a		$C_{19}H_{22}FNO$ m/z 300 (M+H)
68		$C_{20}H_{25}NO_2$ m/z 312 (M+H)

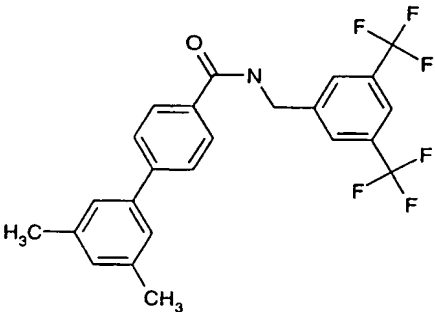
Example #	Structure	Characterization
77		$C_{20}H_{25}NO_2$ m/z 312 (M+H)
78		$C_{19}H_{25}N_3O_3$ m/z 344 (M+H)
79		$C_{18}H_{21}N_5O$ m/z 324 (M+H)
80		$C_{18}H_{22}N_2O$ m/z 283 (M+H)
81		$C_{23}H_{25}NO$ m/z 332 (M+H)

Example #	Structure	Characterization
82		$C_{21}H_{21}F_6NO$ m/z 418 (M+H)
83		$C_{21}H_{27}NO$ m/z 310 (M+H)
84		$C_{23}H_{17}F_6NO_2$ m/z 452 (M-H)
85		$C_{22}H_{14}F_7NO$ m/z 442 (M+H)
86		$C_{23}H_{17}F_6NO_2$ m/z 454 (M+H)

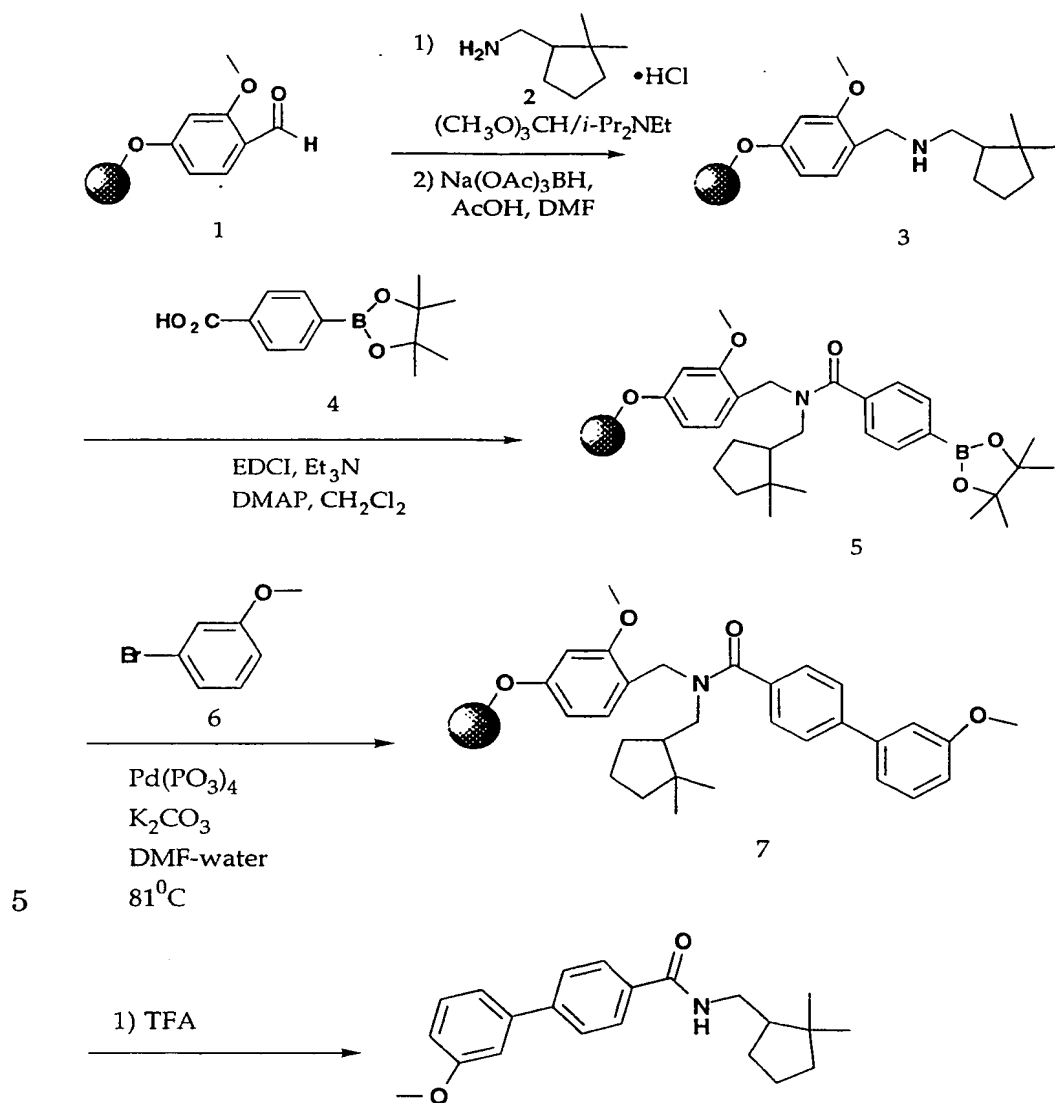
Example #	Structure	Characterization
87		$C_{21}H_{14}F_6N_2O$ m/z 425 (M+H)
88		$C_{26}H_{17}F_6NO$ m/z 474 (M+H)
89		$C_{24}H_{19}F_6NO$ m/z 452 (M+H)
90		$C_{23}H_{17}F_6NO_2$ m/z 454 (M+H)

Example #	Structure	Characterization
91		$C_{22}H_{14}F_7NO$ m/z 442 (M+H)
92		$C_{23}H_{17}F_6NO_2$ m/z 454 (M+H)
93		$C_{23}H_{17}F_6NO_2$ m/z 454 (M+H)
94		$C_{22}H_{17}F_6N_3O_3$ m/z 486 (M+H)

Example #	Structure	Characterization
95		$C_{21}H_{14}F_6N_2O$ m/z 425 (M+H)
96		$C_{26}H_{17}F_6NO$ m/z 472(M-H)
97		$C_{24}H_{13}F_{12}NO$ m/z 558 (M-H)

Example #	Structure	Characterization
98		$C_{24}H_{19}F_6NO$ m/z 452 (M+H)

Example 99



A. Resin 1

10

The title resin was prepared according to Example 68, part A.

B. Resin 3

5.0 g of resin 1 (loading = 1.08 mmol/g, 5.4 mmol) was weighed into a glass SPOS reaction vessel. The resin was swelled with dry DMF (25 ml) which was drained after 10 min. Dry DMF (20 ml) was then added followed by amine 2 (2,2-dimethylmethylaminocyclopentane hydrochloride salt 1.33g, 8.12 mmol) and N,N-diisopropylethylamine (4.7 ml, 27 mmol). The reaction vessel was agitated on a wrist-action shaker for 10 min. and trimethyl orthoformate (7 ml) was added. After 10 min. of agitating for 14 h, the reaction was drained and 20 ml of dry DMF was added. The reaction was agitated for approximately 1 min. and drained. 25 ml of dry DMF was added followed by sodium triacetoxyborohydride (3.40 g, 16.0 mmol) and acetic acid (1.0 ml, 1.6 mmol). After 6 h of agitation, the reaction was drained and rinsed with DMF (3 x 30 ml), DMF-water (3 x 30 ml), DMF (3 x 30 ml), dichloromethane (3 x 30 ml), methanol (3 x 30 ml) and THF (3 x 30 ml). The resin was used in the next step without characterization.

C. Boronate Ester 4

20

To a stirred mixture of 3-carboxyboronic acid (1.65g, 10.0 mmol) in toluene (60 mL) was added pinacol (1.24 g, 10.5 mmol). The reaction flask was fitted with a Dean-Stark trap and a reflux condenser and immersed in an oil bath. The bath was heated to 160°C. After 20 hours, TLC indicated consumption of starting material. The reaction was allowed to cool. The solvent was removed *in vacuo* and the residue purified by flash column chromatography (silica, methanol-dichloromethane, 1:19). The appropriate fractions were collected to give the title compound (2.3 g, 93%).

30

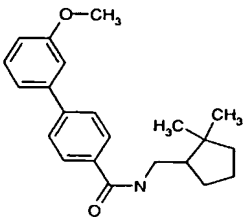
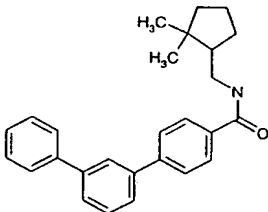
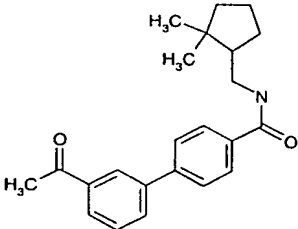
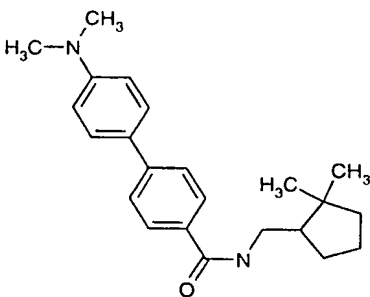
D. Resin 5

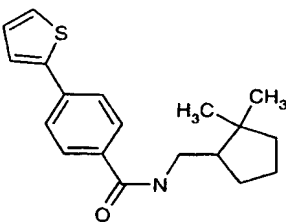
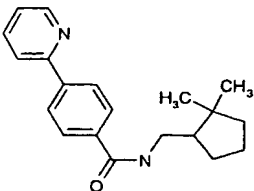
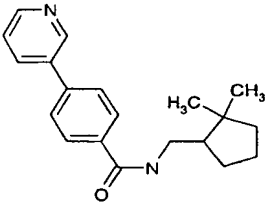
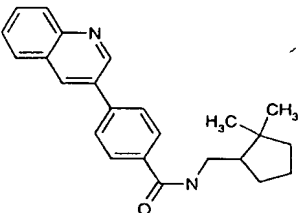
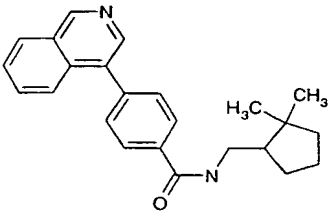
Resin 3 (5.6 g, loading = 0.97 mmol/g, 5.4 mmol) was transferred into a glass SPOS reaction vessel. The resin was swelled with 20 mL of dry dichloromethane. After 10 minutes the solvent was drained away and 5 mL of dry dichloromethane was added. The title 4 compound (1.75 g, 7.06 mmol) was added followed by 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (2.44 g, 7.01 mmol), triethylamine (3.75 mL, 26.9 mmol) and N,N-dimethylaminopyridine (50 mg, cat.). The reaction was agitated on a wrist-action shaker for 16 hour. The solvent was drained from the resin which was then washed with dichloromethane (3 x 40 mL), THF (3 x 40 mL), methanol (3 x 40 mL) and THF (3 x 40 mL).

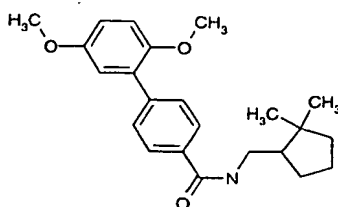
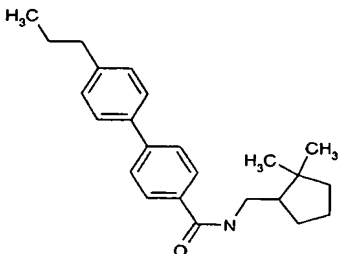
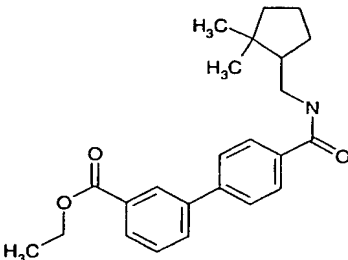
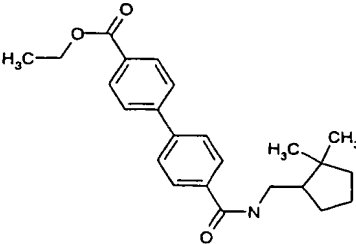
E. Title compound of Example 99

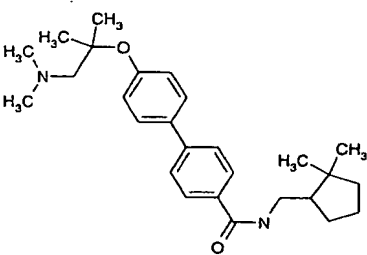
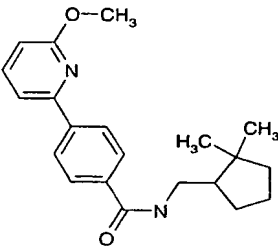
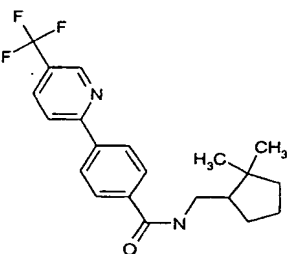
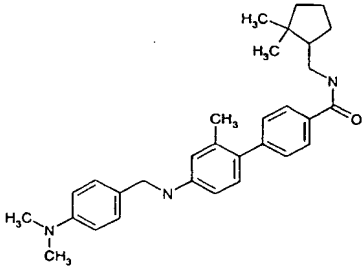
Resin 5 (200 mg, 0.158 mmol) was transferred into a glass reaction tube. DMF-water (9:1, 3 mL) was added followed by 3-bromoanisole (0.50 mL, 0.40 mmol), potassium carbonate (100 mg, 0.72 mmol) and tetrakis(triphenyl-phosphine)palladium-(0) (20 mg, 0.017 mmol). The reaction was sealed and agitated at 250 rpm on an orbital shaker. The reaction was heated to 81°C over a one hour period. After 20 hours, the reaction was allowed to cool to room temperature. The solvent was drained and the resin was washed with DMF (3 x 5 mL), DMF-water (3 x 5 mL), DMF (3 x 5 mL), dichloromethane (3 x 5 mL), THF (3 x 5 mL), methanol (3 x 5 mL), and dichloromethane (3 x 5 mL). Dichloromethane (0.5 mL) was added followed by TFA (3 mL). After 1 hour, the product was collected into a tared test-tube. The solvent was removed from the cleavage product *in vacuo*. The product was reconstituted with 3 mL of dichloromethane and the solvent removed *in vacuo* to provide the title compound (24.1 mg).

Using the method described in Example 99, the following compounds were prepared.

Example #	Structure	Characterization
100		$C_{22}H_{27}NO_2$ m/z 338 (M+H)
101		$C_{27}H_{29}NO$ m/z 384 (M+H)
102		$C_{23}H_{27}NO_2$ m/z 350 (M+H)
103		$C_{23}H_{30}N_2O$ m/z 351 (M+H)

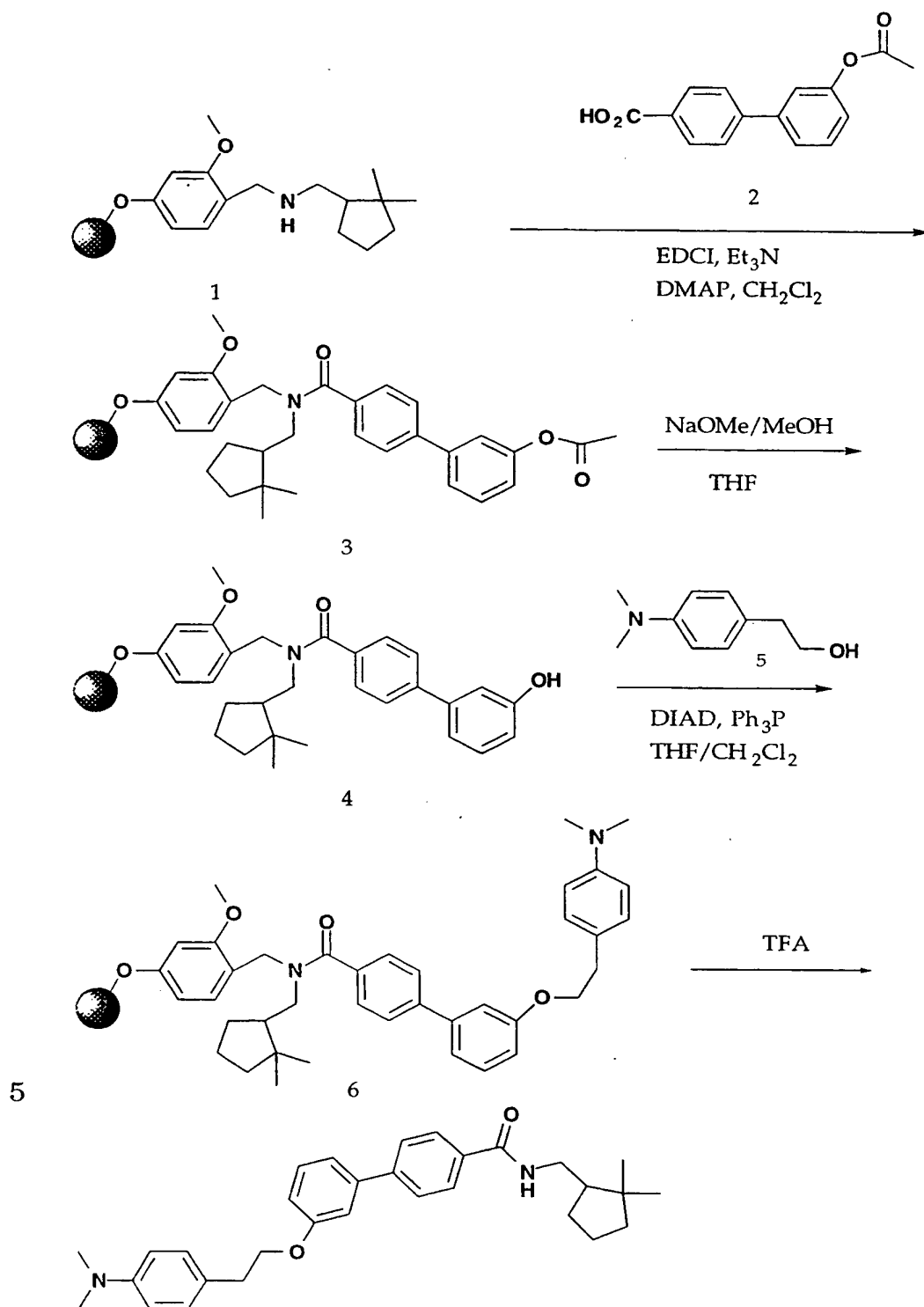
Example #	Structure	Characterization
104	 <chem>CC1(C)CCCC1CN(C(=O)c2ccc(cc2)c3ccccc3S)c4ccccc4</chem>	$C_{19}H_{23}NOS$ m/z 314 (M+H)
105	 <chem>CC1(C)CCCC1CN(C(=O)c2ccc(cc2)c3ccccn3)c4ccccc4</chem>	$C_{20}H_{24}N_2O$ m/z 309 (M+H)
106	 <chem>CC1(C)CCCC1CN(C(=O)c2ccc(cc2)c3ccccn3)c4ccccc4</chem>	$C_{20}H_{24}N_2O$ m/z 309 (M+H)
107	 <chem>CC1(C)CCCC1CN(C(=O)c2ccc(cc2)c3ccc4ccccc4n3)c5ccccc5</chem>	$C_{24}H_{26}N_2O$ m/z 359 (M+H)
108	 <chem>CC1(C)CCCC1CN(C(=O)c2ccc(cc2)c3ccc4ccccc4n3)c5ccccc5</chem>	$C_{24}H_{26}N_2O$ m/z 359 (M+H)

Example #	Structure	Characterization
109		$C_{23}H_{29}NO_3$ m/z 368 (M+H)
110		$C_{24}H_{31}NO$ m/z 350 (M+H)
111		$C_{24}H_{29}NO_3$ m/z 380 (M+H)
112		$C_{24}H_{29}NO_3$ m/z 380 (M+H)

Example #	Structure	Characterization
113		$C_{27}H_{38}N_2O_2$ m/z 423 (M+H)
114		$C_{21}H_{26}N_2O_2$ m/z 339 (M+H)
115		$C_{21}H_{23}F_3N_2O$ m/z 377 (M+H)
116		$C_{31}H_{39}N_3O$ m/z 470 (M+H)

Example #	Structure	Characterization
117		$C_{22}H_{26}N_2O_3$ m/z 367 (M+H)
118		$C_{21}H_{25}N_3O_2$ m/z 352 (M+H)

Example 120



A. Compound 2

A solution of 3-methoxybenzene boronic acid (4.0g, 26.3 mmol) and
5 benzyl-4-bromobenzoate (7.66g, 26.3 mmol) in toluene (50 mL) was
degassed by bubbling nitrogen through for 15 minutes. A solution of
potassium carbonate in water (2 M, 13.1 mL, 26.2 mmol) was added
followed by the addition of tetrakis(triphenylphosphine)palladium (0)
(0.66 g, 0.57 mmol). The reaction flask was fitted with a reflux condenser
10 and immersed in an oil bath which was heated to 150°C. After stirring
under reflux for 14 hours, the reaction was allowed to cool and poured
into an EtOAc (500 mL) - water (200 mL) mixture. The layers were
separated and the organic layer washed with brine, stirred over
magnesium sulfate/charcoal for 15 minutes, filtered through Celite and
15 concentrated *in vacuo* to give 7.5 g of a white solid which was dissolved
in ethanol and added to a pressure flask. Palladium-hydroxide on
carbon (10%, 750 mg) was added. The flask was placed on a Parr shaker
and agitated under a hydrogen atmosphere (63 psi) for 4 hours. At the
end of this time, nitrogen was bubbled through the reaction mixture for
20 15 minutes. The reaction mixture was filtered through a pad of Celite
and concentrated in vacuum to afford 5.32 g of a white solid. Dry
dichloromethane (100 mL) was added to the flask containing the above
material. The flask was cooled to -78°C in a dry ice - acetone bath and a
solution of boron tribromide in dichloromethane (1 M, 57 mL, 57 mmol)
25 was added via syringe over a period of 20 minutes. The reaction was
stirred at -78°C for 2 hours. At this time the dry ice-acetone bath was
replaced with an ice-water bath and the reaction was allowed to warm to
4°C over the period of 1 hour. At this time, the reaction was placed in the
dry ice-acetone bath. After 10 minutes, methanol (10 mL) was added via
30 syringe over a 2 minute period. The reaction was poured into a
dichloromethane (500 mL) - water (100 mL) mixture. The layers were
separated and the aqueous layer was extracted with dichloromethane (5

x 200 mL). The dichloromethane extracts were combined, dried over magnesium sulfate and concentrated *in vacuo*. The resulting material was purified by flash column chromatography (silica, 9:1 - dichloromethane : methanol). The appropriate fractions were pooled
5 and concentrated *in vacuo* to afford 2.99 g of a white solid. The above material was suspended in acetic anhydride and 3 drops of an acetic acid:sulfuric acid (1:1) mixture were added. The reaction flask was immersed in an oil bath which was heated to 160°C for 1.5 hours. At the end of this time, the reaction was poured over an ice (50 g)/water (50 mL)
10 mixture. The flask containing this mixture was heated at 55°C with stirring for 1 hour. Over this period a white precipitate appeared in the flask. The flask was cooled in an ice-water bath for 1/2 hour and the precipitate was collected by vacuum filtration. This procedure yielded 3.3 g of 2.

B. Resin 3

Resin 1 (the title A resin of example 99, 5.6 g, loading = 0.97 mmol/g, 5.4 mmol) was transferred into a glass SPOS reaction vessel.
20 The resin was swelled with 20 mL of dry dichloromethane. After 10 minutes the solvent was drained away and 25 mL of dry dichloromethane was added. Acid 2 (2.07 g, 10.5 mmol) was added followed by 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (2.82 g, 10.5 mmol), triethylamine (1.8 mL, 12.9 mmol)
25 and N,N-dimethylaminopyridine (20 mg, cat.). The reaction was agitated on a wrist-action shaker for 16 hours. The solvent was drained from the resin which was then washed with dichloromethane (3 x 40 mL), THF (3 x 40 mL), methanol (3 x 40 mL) and THF (3 x 40 mL).

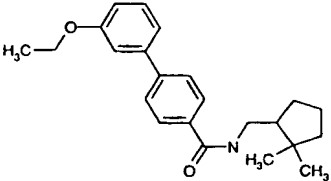
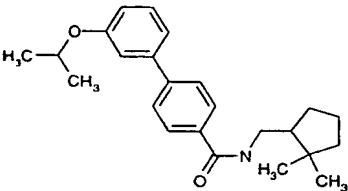
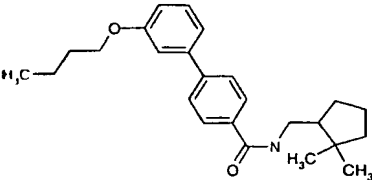
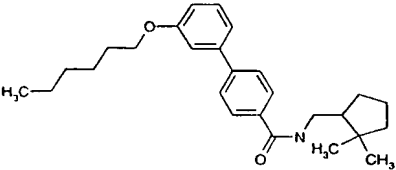
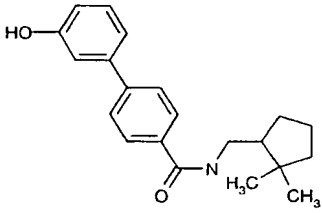
C. Resin 4

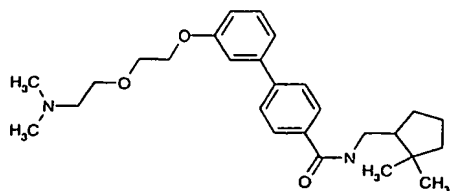
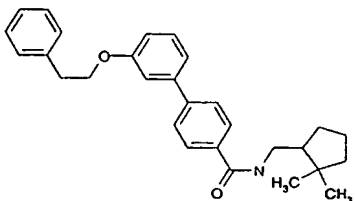
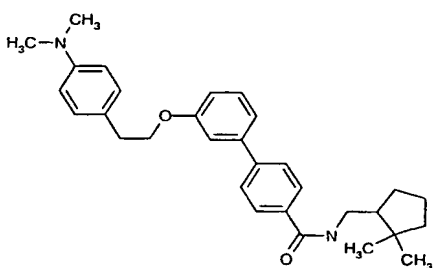
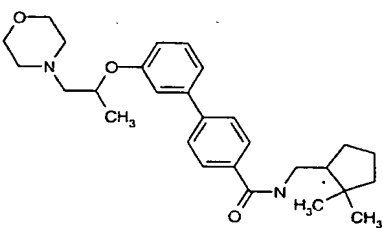
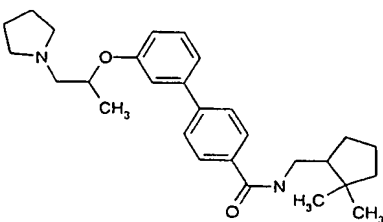
6.0 g of resin 3 (loading = 0.78 mmol/g, 4.7 mmol) was transferred into a glass SPOS reaction vessel. The resin was swelled with 50 mL of THF and 20 mL of a solution of sodium methoxide in methanol (0.5 M, 10 mmol) was added. The reaction was agitated for 20 hours and drained. The resin was washed with THF (40 mL) which was drained. A solution of acetic acid in THF (1:9) was then added. After agitating for 1/2 hour, the reaction was drained and the resin was then washed with dichloromethane (3 x 40 mL), THF (3 x 40 mL), methanol (3 x 40 mL) and THF (3 x 40 mL).

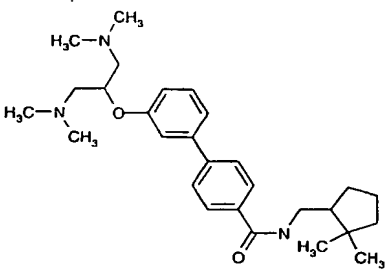
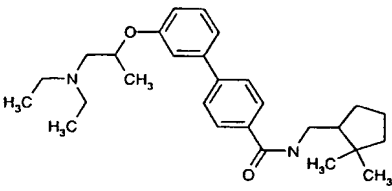
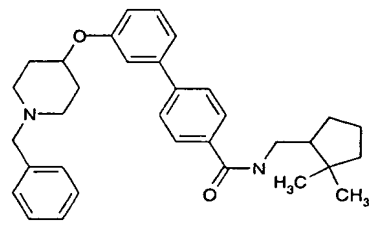
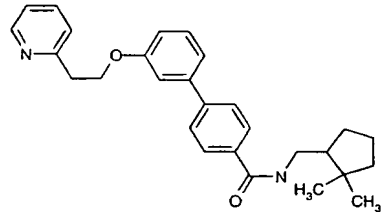
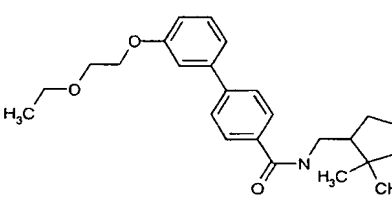
D. The title compound of Example 120

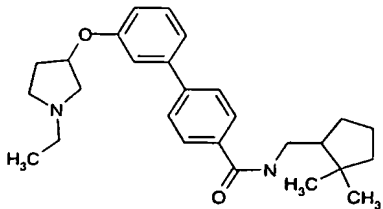
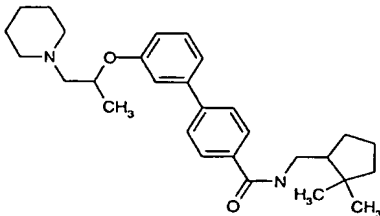
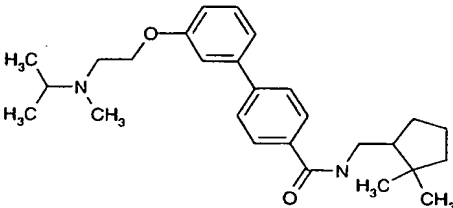
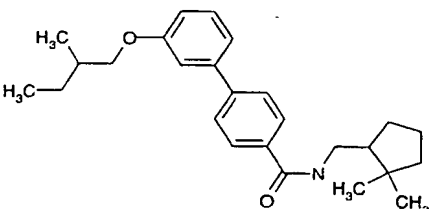
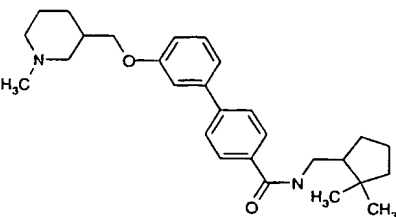
Resin 4 (200 mg, 0.164 mmol) was transferred into a polypropylene reaction tube which was fitted into a solid-phase reactor. The resin was swelled by the addition of 2.0 mL of dichloromethane. To the resin was added 4-(dimethylamino)-phenethyl alcohol (136 mg, 0.82 mmol), 1 mL of a solution of triphenylphosphine in THF (0.82 M, 0.82 mmol) and 1 mL of a solution of diisopropyl azodicarboxylate in THF (0.82 M, 0.82 mmol). The reaction block was agitated at 275 rpm for 20 hours. At the end of this period the solvent was drained and the resin was washed with DMF (3 x 5 mL), DMF-water (3 x 5 mL), DMF (3 x 5 mL), dichloromethane (3 x 5 mL), THF (3 x 5 mL), methanol (3 x 5 mL), and dichloromethane (3 x 5 mL). Dichloromethane (0.5 mL) was added followed by trifluoroacetic acid (3 mL). After 1 hour, the product was collected into a tared test-tube. The solvent was removed from the cleavage product *in vacuo*. The product was reconstituted with 3 mL of dichloromethane and the solvent removed *in vacuo* to provide the title compound (30.3 mg).

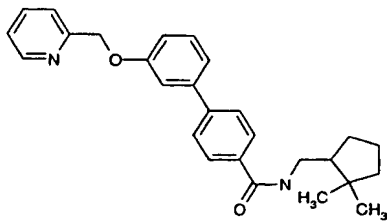
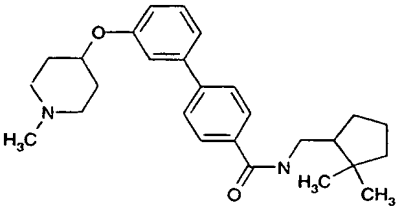
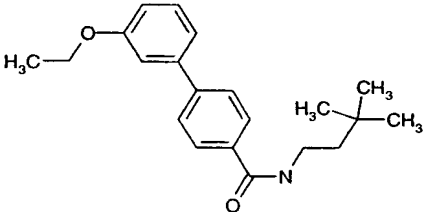
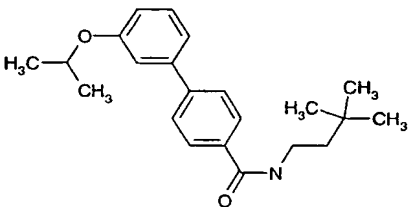
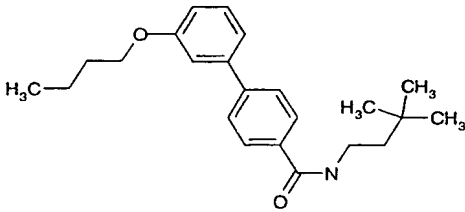
Using the procedure described in example 120, the following compounds were prepared.

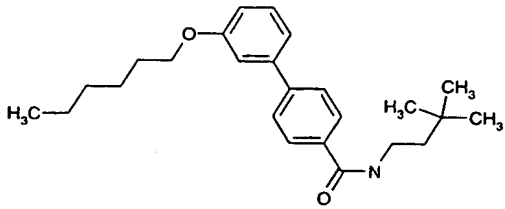
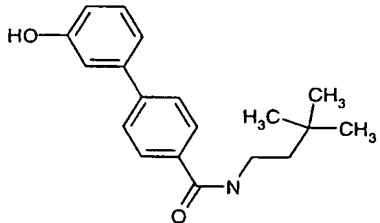
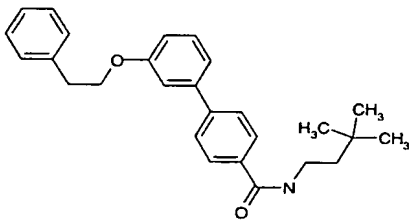
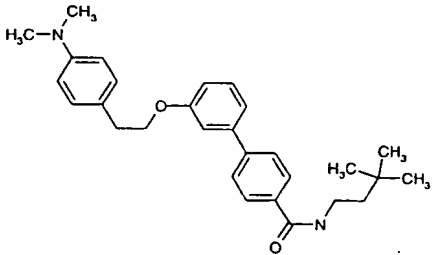
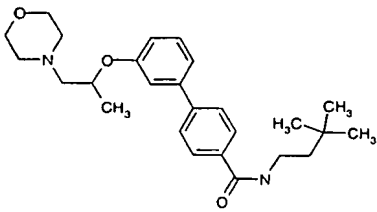
Example #	Structure	Characterization
121		$C_{23}H_{29}NO_2$ m/z 352 (M+H)
122		$C_{24}H_{31}NO_2$ m/z 366 (M+H)
123		$C_{25}H_{33}NO_2$ m/z 380 (M+H)
124		$C_{27}H_{37}NO_2$ m/z 408 (M+H)
125		$C_{21}H_{25}NO_2$ m/z 324 (M+H)

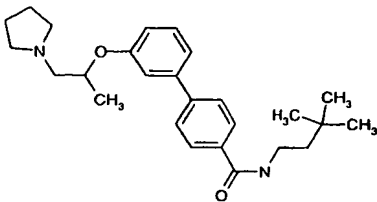
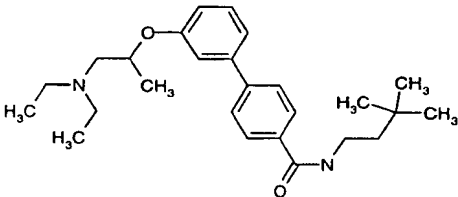
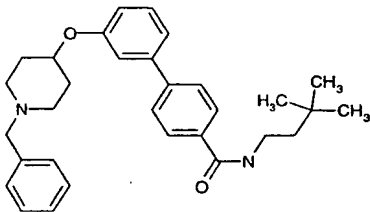
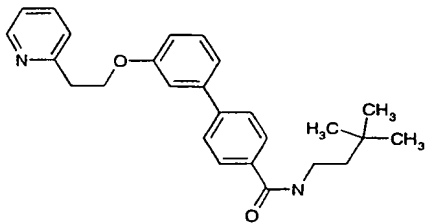
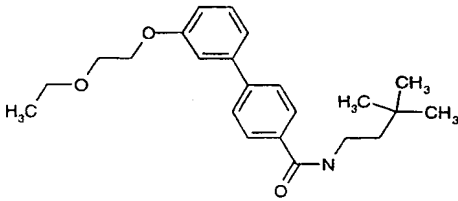
Example #	Structure	Characterization
126		$C_{27}H_{38}N_2O_3$ m/z 439 (M+H)
127		$C_{29}H_{33}NO_2$ m/z 428 (M+H)
120		$C_{31}H_{38}N_2O_2$ m/z 471 (M+H)
128		$C_{28}H_{38}N_2O_3$ m/z 451 (M+H)
129		$C_{28}H_{38}N_2O_2$ m/z 435 (M+H)

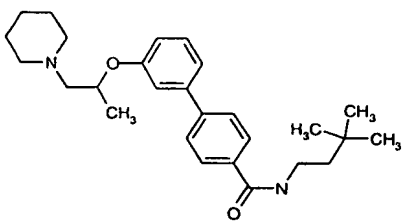
Example #	Structure	Characterization
130		$C_{28}H_{41}N_3O_2$ m/z 452 (M+H)
131		$C_{28}H_{40}N_2O_2$ m/z 437 (M+H)
132		$C_{33}H_{40}N_2O_2$ m/z 497 (M+H)
133		$C_{28}H_{32}N_2O_2$ m/z 429 (M+H)
134		$C_{25}H_{33}NO_3$ m/z 396 (M+H)

Example #	Structure	Characterization
135		$C_{27}H_{36}N_2O_2$ m/z 421 (M+H)
136		$C_{29}H_{40}N_2O_2$ m/z 449 (M+H)
137		$C_{27}H_{38}N_2O_2$ m/z 423 (M+H)
138		$C_{26}H_{35}NO_2$ m/z 394 (M+H)
139		$C_{28}H_{38}N_2O_2$ m/z 435 (M+H)

Example #	Structure	Characterization
140		$C_{27}H_{30}N_2O_2$ m/z 415 (M+H)
141		$C_{27}H_{36}N_2O_2$ m/z 421 (M+H)
142		$C_{21}H_{27}NO_2$ m/z 326 (M+H)
143		$C_{22}H_{29}NO_2$ m/z 340 (M+H)
144		$C_{23}H_{31}NO_2$ m/z 354 (M+H)

Example #	Structure	Characterization
145		$C_{25}H_{35}NO_2$ m/z 382 (M+H)
146		$C_{21}H_{23}NO_2$ m/z 298 (M+H)
147		$C_{27}H_{31}NO_2$ m/z 402 (M+H)
148		$C_{29}H_{36}N_2O_2$ m/z 445 (M+H)
149		$C_{26}H_{36}N_2O_3$ m/z 425 (M+H)

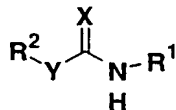
Example #	Structure	Characterization
150		$C_{26}H_{36}N_2O_2$ m/z 409 (M+H)
151		$C_{26}H_{38}N_2O_2$ m/z 411 (M+H)
152		$C_{31}H_{38}N_2O_2$ m/z 471 (M+H)
153		$C_{26}H_{30}N_2O_2$ m/z 403 (M+H)
154		$C_{23}H_{31}NO_3$ m/z 370 (M+H)

Example #	Structure	Characterization
155	 <chem>CC(C)(C)CN(C(=O)c1ccc(cc1)-c2ccc(OC)cc2N3CCCCC3)C4=CC=CC=C4</chem>	$C_{27}H_{38}N_2O_2$ m/z 423 (M+H)

What is Claimed is:

1. A method of treating cardiac arrhythmia which comprises
 5 administering to a mammal in need thereof an effective amount of a compound of the formula

I



where

- 10 X is oxygen, sulfur, -NH, -NR¹, -N-CN, -N-OR¹ or
 -N-NO₂;

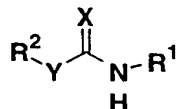
Y is a single bond, -C=C-, or -NH;

R¹ is alkyl, alkenyl, alkynyl, aryl, cycloalkyl, heterocyclo, or
 (heterocyclo)alkyl; and

- 15 R² is aryl or heterocyclo.

2. A compound of the formula

I



- 20 where

X is oxygen, sulfur, -NH, -NR¹, -N-CN, -N-OR¹ or
 -N-NO₂;

Y is a single bond, -C=C-, or -NH;

- R¹ is alkyl, alkenyl, alkynyl, aryl, cycloalkyl, heterocyclo, or
 25 (heterocyclo)alkyl; and

R² is aryl or heterocyclo.

3. A compound as recited in Claim 2 wherein

X is oxygen or N-CN;

Y is a single bond or NH; and

5 R¹ is alkyl, cycloalkyl, (aryl)alkyl, (cycloalkyl)alkyl, or (substituted-amino)alkyl.

4. A compound as recited in Claim 2 wherein

X is oxygen;

Y is a single bond or NH; and

10 R¹ is alkyl, cycloalkyl, (aryl)alkyl, (cycloalkyl)alkyl, or (substituted-amino)alkyl.

5. A compound as recited in Claim 2 wherein

X is N-CN;

15 Y is a single bond or NH; and

R¹ is alkyl, cycloalkyl, (aryl)alkyl, (cycloalkyl)alkyl, or (substituted-amino)alkyl.

6. A compound as recited in Claim 2 wherein

20 X is oxygen or N-CN;

Y is NH; and

R¹ is alkyl, cycloalkyl, (aryl)alkyl, (cycloalkyl)alkyl, or (substituted-amino)alkyl.

25 7. A compound as recited in Claim 2 wherein

X is oxygen;

Y is a single bond; and

30 R¹ is alkyl, cycloalkyl, (aryl)alkyl, (cycloalkyl)alkyl, or (substituted-amino)alkyl.

8. A compound as recited in Claim 2 wherein
X is N-CN
Y is a single bond; and
R¹ is alkyl, cycloalkyl, (aryl)alkyl, (cycloalkyl)alkyl, or (substituted-
5 amino)alkyl.
9. A compound as recited in Claim 2 wherein
X is oxygen or N-CN;
Y is single bond or NH; and
10 R¹ is alkyl.
10. A compound as recited in Claim 2 wherein
X is oxygen or N-CN;
Y is single bond or NH and
15 R¹ is cycloalkyl.
11. A compound as recited in Claim 2 wherein
X is N-CN;
Y is single bond or NH; and
20 R¹ is (aryl)alkyl.
12. A compound as recited in Claim 2 wherein
X is N-CN;
Y is single bond or NH; and
25 R¹ is (cycloalkyl)alkyl.
13. A compound as recited in Claim 2 wherein
X is N-CN;
Y is single bond or NH and
30 R¹ is (substituted-amino)alkyl.

14. A compound as recited in Claim 2 which is:

(+)-N-[(2,2-dimethylcyclopentyl)methyl]-4-hexyloxybenzamide and
 (-)-N-[(2,2-dimethylcyclopentyl)-methyl]-4-hexyloxybenzamide;

N-(3,3-dimethylbutyl)-4-(1H-indol-1-yl)benzamide;

5 4-(3-butyl-1,2,4-oxadiazol-5-yl)-N-(3,3-dimethylbutyl)-benzamide;

N-(3,3-dimethylbutyl)-2-(hexyloxy)-5-pyridine carboxamide;

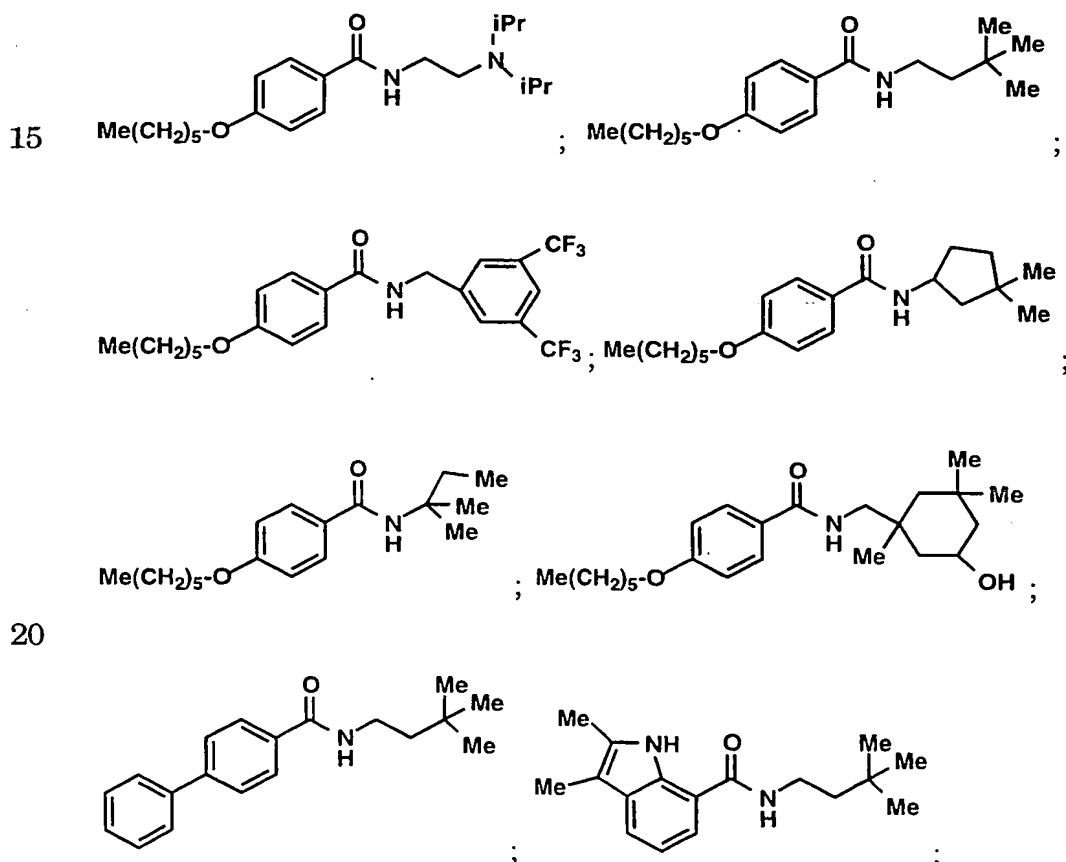
N-(3,3-dimethylbutyl)-1-hexyl-1,2-dihydro-2-oxo-5-pyridinecarboxamide;

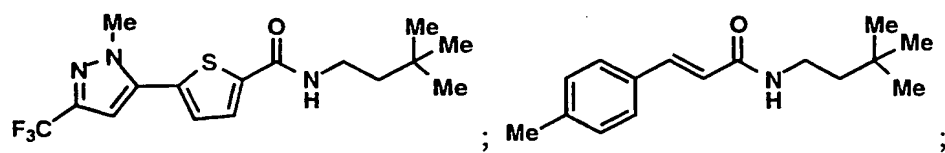
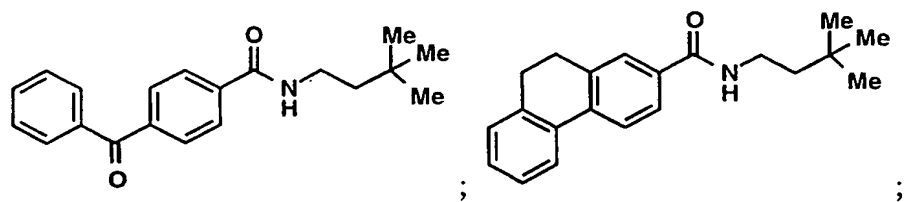
10 N-cyano-N'-(3,3-dimethylbutyl)-6-(hexyloxy)benzene-carboximidamide;

N-(3,3-dimethylbutyl)-N'-[4-(hexyloxy)phenyl]urea;

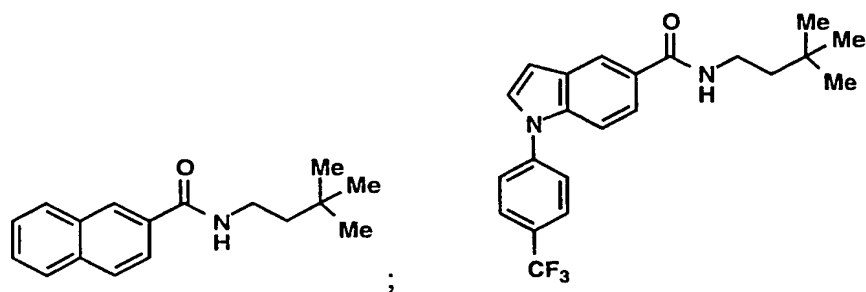
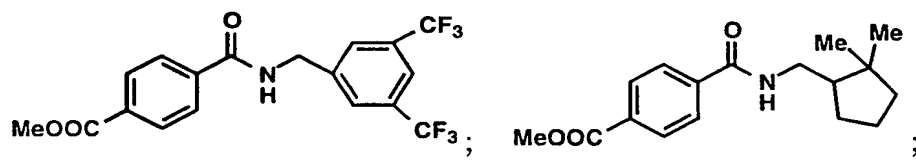
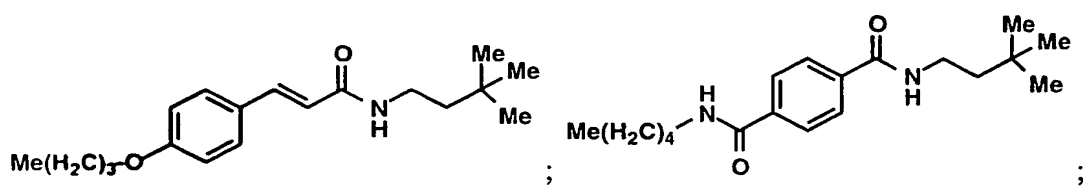
N-(3,3-dimethylbutyl)-N'-[4-(hexyloxy)phenyl]urea;

or a compound of claim 2 having the structure;

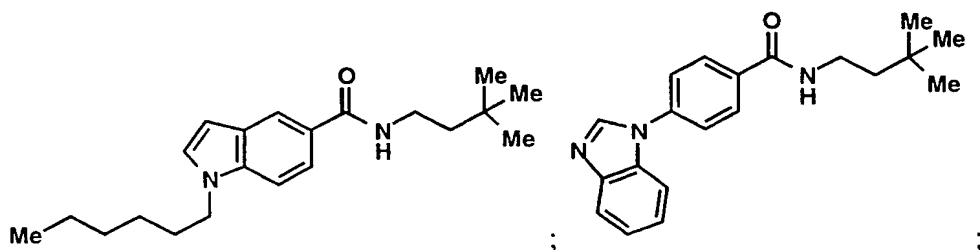


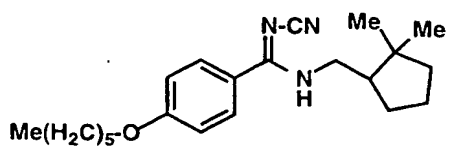
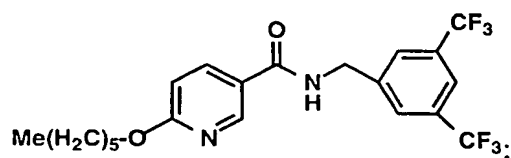
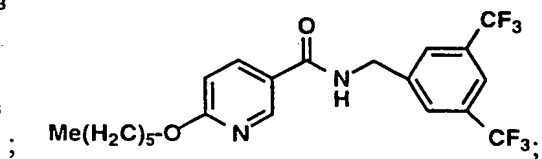
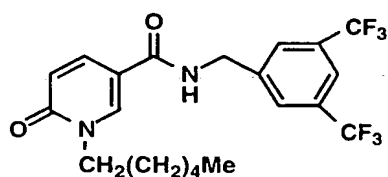
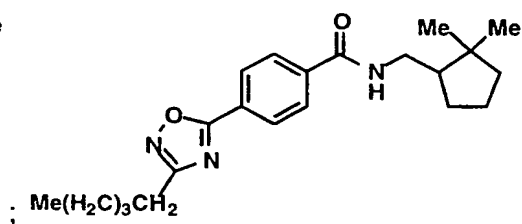
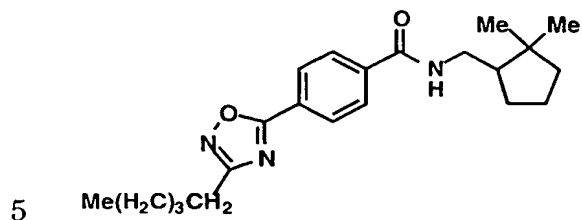
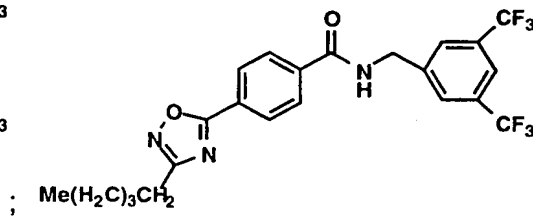
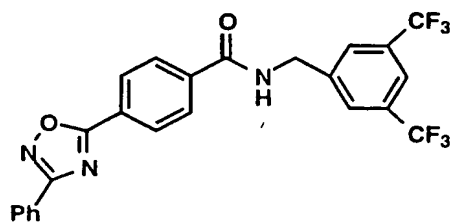
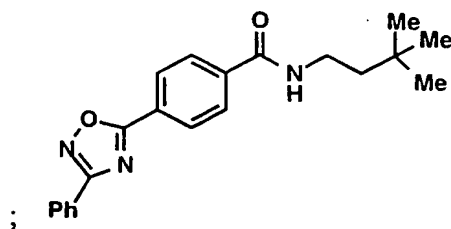
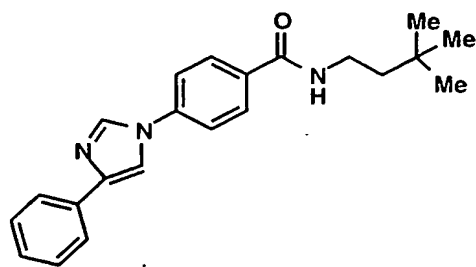


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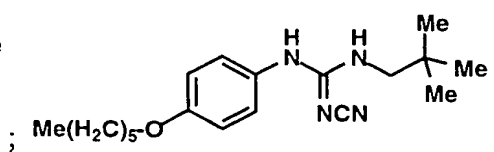
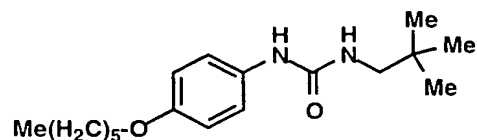


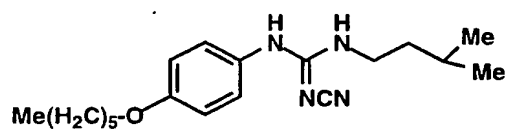
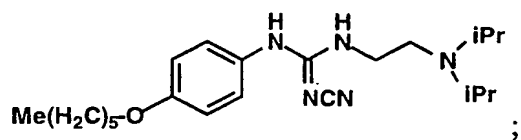
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; or a pharmaceutically acceptable salt

5 thereof.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US98/02364

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : C07D 209/04, 231/10, 231/56, 235/08

US CL : 548/304.4, 357.5, 469

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 548/304.4, 357.5, 469

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS ONLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,098,924 (POSS) 24 March 1992, see the entire document, especially pages 66-88.	1-14

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A* document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means	
P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

13 APRIL 1998

Date of mailing of the international search report

08 JUN 1998

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

TAOFIQ A. SOLOLA

Telephone No. (703) 308-1235

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